

GenCore version 5.1.3
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SUMMARIES

OM protein - nucleic search, using frame_plus_p2n model

Run on: October 19, 2002, 07:51:16 : Search time 199 Seconds

(without alignments)
3951.492 Million cell updates/sec

Title: US-09-807-459-2

Perfect score: 2359

Sequence: 1 MAPSDSVGDVTKTLAASES.....DPKALIRKVSSTEADNLEK 458

Scoring table:

BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

-MODL=frame+ p2n.model -DEV=xlh
-O=/cgn2_1/USPRO/spool/US09807459/runat_18102002_141111-28571/app-query.fasta.1.647
-DB=N_Geneseq_032802 -QFMT=fastap -SUFFIX=ring -MINMATCH=0.1 -LOOPEXT=0
-LOOPEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=blomsum62 -TRANS=human40.cdi
-LIST=45 -DOCALLIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=40
-MODE=LOCAL -OUTFMT=prc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09807459 -GCN_1_1_48 -runat_18102002_141111-28571 -NCPU=6 -ICPU=3
-NO_XLPRY -NO_MAP -LARGEQUERY -NEG SCORES=0 -WAIT -LONGLOG -DEV=TIMEOUT=120
-WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6 -Fgapext=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : N_Geneseq_032802:*

1: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA1980.DAT:*
2: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA1981.DAT:*
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22: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2001A.DAT:*
23: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2001B.DAT:*
24: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

Result No.	Score	Query Match	Length	DB ID	Description
1	2359	100.0	1828	22 AAF59961	Babesia caballi me
2	996.5	42.2	1542	24 AAO47074	21B4/rhoptry genes
3	827.5	35.1	1962	16 AAO90252	Babesia merzoiite
4	826.5	35.0	1990	14 AAO33064	Encodes Babesia bo
5	826.5	35.0	1990	17 AAT18995	Babesia merzoiite
6	770.5	32.7	1371	14 AAO47076	B. canis 21B4/rhop
7	759.5	32.2	1491	14 AAO47075	21B4/rhoptry anti
8	161	6.8	282	13 AAO26065	21B4 gene clone pr
9	147	6.2	1664976	19 AAV71209	Methanococcus jam
10	138	5.8	170	13 AAO26066	21B4 gene clone pr
11	134	5.7	26776	20 AAX20254	Borrelia burgdorfe
12	122.5	5.2	111309	20 AAX20250	Borrelia burgdorfe
13	122.5	5.2	910715	20 AAX20248	Borrelia burgdorfe
14	121	5.1	11165	21 AAI15186	DNA encoding Esche
15	120	5.1	3883	20 AAV71915	S. cerevisiae C1N8
16	120	5.1	3884	22 AAH78010	Nucleotide sequenc
17	118.5	5.0	4766	15 AAO70102	Malaria P1EMP3 ep
18	118.5	5.0	7326	21 AAA70095	Plasmodium falcipa
19	117	5.0	33303	21 AAA81514	N. meningitidis pa
20	117	5.0	349980	21 AAF21610	Neisseria meningit
21	115	4.9	6637	23 AAS71042	DNA encoding novel
22	113.5	4.8	4443	23 ABL04821	Drosophila melanog
23	113.5	4.8	10478	20 AAV84691	Arabidopsis ESD4 (
24	113	4.8	4282	23 AAS76165	DNA encoding novel
25	113	4.8	11087	23 AAS7637	DNA encoding novel
26	112.5	4.8	2375	22 AAK68993	Human immune/haema
27	112.5	4.8	42738	22 AAK68992	Human immune/haema
28	111	4.7	1422	20 AAX35720	cDNA encoding a pr
29	111	4.7	1613	20 AAX35721	K. lactis origin o
30	111	4.7	3278	18 AAT73285	Kluyveromyces lact
31	111	4.7	3278	18 AAT73285	DNA segment 4 codi
32	110	4.6	1777	11 AAO06842	DNA encoding novel
33	109.5	4.6	1816	23 AAS76418	DNA encoding novel
34	109.5	4.6	2277	23 AAS69641	DNA encoding novel
35	109.5	4.6	2277	23 AAS71279	DNA encoding novel
36	109.5	4.6	2277	23 AAS74306	DNA encoding novel
37	109.5	4.6	2277	23 AAS74426	DNA encoding novel
38	109.5	4.6	2277	23 AAS74599	DNA encoding novel
39	109.5	4.6	2277	23 AAS79126	DNA encoding novel
40	109.5	4.6	2394	23 AAS84093	DNA encoding novel
41	109.5	4.6	3569	23 AAS78798	DNA encoding novel
42	109.5	4.6	1664976	19 AAV21209	Methanococcus jam
43	109	4.6	1572	20 AAX61708	B. burgdorferi ant
44	109	4.6	2746	23 AAS82185	DNA encoding novel
45	109	4.6	3165	23 AAS78781	DNA encoding novel

ALIGNMENTS

RESULT 1
AAF59961
AAF59961 standard: cDNA, 1828 BP.

22-MAY-2001 (first entry)

Babesia caballi merzoiite 48 kd rhoptry protein-encoding cDNA.
Merzoiite protein: 48 kd rhoptry protein; antigen; antibody;
recombinant production; diagnosis; equine babesiosis;
parasitic infection; veterinary; ss.

Babesia caballi.

WO200112813-A1.

PD 22-FEB-2001.
XX
PF 13-AUG-1999; 99MO-JP04386.
XX
PR 13-AUG-1999; 99MO-JP04386.
XX
PA (KAGA) CHERO-THERAPEUTIC RES INST.
PI (MIKA) MIKAMI T.
XX
PI Mkami T, Ikadei H, Igarashi I, Suzuki N, Nagasawa H, Fujisaki K;
DR WPI: 2001-202867/20.
DR P-PSDB: AAB60669.
XX
PT Gene encoding merozoite protein of Babesia caballi for diagnosis of
XX equine babesiosis caused by this organism -
PS Claim 3; Page 19-22; 27pp: Japanese.
XX
CC The invention relates to a 48 kD merozoite rhoptry protein from Babesia
CC caballi (AAB60669) and cDNA encoding it (AAF59961). The invention also
CC relates to phase vectors containing a nucleic acid encoding the
CC merozoite protein, a method for the recombinant production of the
CC protein, an antibody against the protein, and a method for the diagnosis
CC of equine babesiosis from horse blood samples by using the antibody to
CC detect Babesia caballi merozoites, or by using the 48 kD protein as an
CC antigen to detect anti-Babesia caballi antibodies. The 48 kD merozoite
CC protein, or an antibody specific for the protein may be used for the
CC diagnosis of equine babesiosis caused by Babesia caballi. The present
CC sequence represents cDNA encoding the Babesia caballi merozoite 48 kD
CC rhoptry protein.
XX
SQ Sequence 1828 BP; 523 A; 412 C; 460 G; 433 T; 0 other:
Alignment Scores:
Pred. No.: 1.46e-222 Length: 1828
Score: 2359.00 Matches: 458
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: Gaps: 0
US-09-807-459-2 (1-458) x AAF59961 (1-1828)
QY 1 MetAlaProSerAspSerValIGlyAspValThrLysThrLeuAlaAlaSerGluSer 20
DB 39 ATGGCTCCAGCGACTGTGGGAGCTGACTAGACCTTATGGCTCCAGCGAAAGT 98
QY 21 ValAspSerAlaAlaAsnAlaTyrMetCLeuAsnSerAspMetSerAspTyrLeuSerAla 40
DB 99 GTGACTCAGCTGCCAATGCTATATGATCAACAGTACATGACATGACGATTACTTGTGGCT 158
QY 41 ValSerAspAsnPheAlaGluArgIleCysSerGlnValProLysGlySerAsnCysSer 60
DB 159 GTGTCTACACCTTGGCGAGCGCATTTGCAAGGCTCCCTAAGGGAGAGTAACGTGAGT 218
QY 61 AlaSerValSerAlaTyrMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSer 80
DB 219 GCTTCGCTTAGCGCATGACATGAGCTGCGCTCAACAGAGCTCCGACCTCTCAAAAGT 278
QY 81 LeuLysTyrProLeuGlnAlaLysTyrGlnProLeuThrLeuProAspProTyrGlnLeu 100
DB 279 CTTAAAGTACCTCTTGAAGGCTAAAGTACCAACCCCTGACCCCTTACCACTTGG 338
QY 101 GluAlaAlaPheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSerThrGluLys 120
DB 339 GAGGCCCATTTTACTCTTCAAGAGAGTACCTAATCCGGCAATAGCATGAGAAAG 398
QY 121 ArgPheThrPheAlaArgArgGlyLysAsnHisSerTyrPheHisAspLeuValPhe 140
DB 399 CGGTTCTGATGCGCTTTCAGAGGGGCAAGACACAGTACTTCCACGACTTGTCTTC 458
QY 141 AsnLeuLeuLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsnPheAla 160

DB 459 AATCTGCTGGAGAAAGACGTGACTCGACCGGATGCTACTGACATTTGACCTTGGCG 518
QY 161 SerArgTyrLeuTyrMetAlaThrLeuTyrTyrLysThrTyrTrpAsnValAspGluPhe 180
DB 519 TCCAGTACCTCTTACATGACGCGACCTTTACTACAAAGCATACGAATGTGATGAGTTC 578
QY 181 GlyAlaSerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrGlyLys 200
DB 579 GGTGCTAGCTCTTTAAACAAGTGTCTTACACTGCTGAGGTGTTCGCGCTGGGATCAAG 638
QY 201 ArgAlaLeuLysGlnIleIleArgSerAsnLeuProLeuAspIleGlyThrGluHisSer 220
DB 639 AGGCACCTTAAGCAGATTATTCGCTTAACCTGCCCTTGACATCGGAGACAGAACAGC 698
QY 221 ValSerArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIlePro 240
DB 699 GTCAAGTCCCTGACAGCACATTACGACGATTACAGGATTTACATGATACGACATTCCT 758
QY 241 AlaLeuProLysPheAlaLysArgPheSerLeuMetValGlnArgLeuLeuAlaThr 260
DB 759 GCACCTGCCAAGTTTGGCAACGTTTCTCCTTATGGTACGACAGAGCTGTGGCCACC 818
QY 261 ValAlaGlyTyrValAspThrProTyrTyrLysTyrTyrMetLysLeuLysAsnPhe 280
DB 819 GTGCTGCTTACGTTCGACACCCCGTGGTATAGAAAGTGTACATGAAAGCTGAAACACTT 878
QY 281 MetValAsnArgValPheIleProThrLysPhePheAsnLysGluIleArgGluPro 300
DB 879 ATGGTGAACAGGTTGTTATTCCTACAAAGATTTCTCATTAAGAAATTCGAGACT 938
QY 301 SerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeuPheGluAsnLysIle 320
DB 939 AGTAAGGATTTAAAAAGAAAGTGTCAACCGACACCAAGATTTATTCGAGAAACAAATTT 998
QY 321 GlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspProSerLysAlaLeuLys 340
DB 999 GGGCAGGTTACTGTGCGACTTCTCAATAGGAATTTGCTGACCTAGTAAGGCATTAAAA 1058
QY 341 GluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThrVal 360
DB 1059 GAANAAGTGTCAACGACGCCCAAGATTTATTCAGAACAAATTTGGCAGGCTACTGTG 1118
QY 361 AspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValSerThr 380
DB 1119 GACTTCAATCAATTAACGAATTCGTGACCTAGTAAGCATTAATTAAGAAAGTGTCAACG 1178
QY 381 GlyValaGluAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnAsn 400
DB 1179 GGGCCGAGGATTTATTCGAGAAACAAATTTGGGAGGCTACTGTGACTTCATCAATAAC 1238
QY 401 GluIleArgAspProSerLysAlaLeuIleArgLysValTyrThrGluAlaAspLeu 420
DB 1239 GAATTTGCTGACCTTAGTAAGCATTTAATTAAGAAAGTGTACACCGAGCGCATTTTA 1298
QY 421 PheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnLysGluIleArgAspPro 440
DB 1299 TTTCAGAAACAAATTTGGGAGGCTAGTGTGACTTCATCAATTAAGAAATTCGTGACCT 1358
QY 441 SerLysAlaLeuIleArgLysValSerThrGluAlaAspAsnLeuLeuGluLys 458
DB 1359 AGTAAGCATTAATTAAGAAAGTGTTCACGAGGCGCATTAATTTATTTGAGAAA 1412
RESULT 2
ID AA047074 standard; DNA: 1542 BP.
XX
AC AA047074;
XX
DT 13-JAN-1994 (first entry)
XX
DE 21B4/rhoptry genes 1-4 representative DNA.
XX

Db 1369 GGTGCCCCCATTAAGA-ATCATTTGAGAACTGGACACAGCG 1412
RESULT 3
ID AAQ90252 standard; cDNA: 1962 BP.
XX
AC AAQ90252;
XX
DE 14-NOV-1995 (first entry)
XX
DE Babesia merozoit p58 cDNA.
XX
KM Merozoite; surface protein; antigen; p58; babesiosis; vaccine; ss.
OS Babesia bigemina.
XX
FH Key Location/Qualifiers
FT CDS 186..1628
FT /tag- a
FT sig_peptide 186..248
FT /tag- b
FT mat_peptide 249..1631
FT /tag- c
FT polyA_signal 1884..1889
FT /tag- d
XX
PN US542428-A.
XX
PD 06-JUN-1995.
XX
PF 27-MAR-1987; 87US-0031328.
XX
PR 06-DEC-1991; 91US-0803636.
PR 27-MAR-1987; 87US-0031328.
PR 01-MAR-1991; 91US-0663255.
XX
PA (UNIW) UNIV WASHINGTON STATE.
XX
PI Davis WC, McElwain TF, McGuire TC, Perryman LE;
XX
DR MPI: 1995-214706/28.
DR P-PSDB; AAR77249.
XX
PT Babesia merozoit 45 kD surface protein from B. bigemina - used in
XX vaccines for the prophylaxis of bovine babesiosis.
XX
PS Disclosure: Column 31-34; 30pp: English.
XX
CC Antigenic surface proteins (45, 55 and 58 kDa) were isolated from
CC the intracythrotic merozoit stage of B. bigemina JG-29. The 58
CC kDa surface protein (AAR77249) was characterized, and encoding
CC cDNA (AAQ90252) was isolated from a lambda GEM1 library.
XX
SO Sequence 1962 BP: 506 A; 442 C; 492 G; 522 T; 0 other:
SO
Alignment Scores:
Pred. No.: 1e-71 Length: 1962
Score: 827.50 Matches: 181
Percent Similarity: 59.12% Conservative: 75
Best Local Similarity: 41.80% Mismatches: 140
Query Match: 35.08% Indels: 37
DB: 16 Gaps: 12
US-09-807-459-2 (1-458) x AAQ90252 (1-1962)
OY 4 SerAspSerValGlyAspValThrIysThrLeuAlaIaSerGluSerValAspSer 23
DB 282 GAGAGGGGTGGAGATGTCTCAAGACCTTGCGAAGCCAAATGAGGTTGCAATGCT 341
OY 24 AlaIaAsnAlaTyrMetIleAsnSerAspMetSerAspTyrLeuSerAlaValSerAsp 43
DB 342 GAATGGAAGCAACATCAGGTCAACAAGATATGCAAAAGCATTTGTCTAATGTTAAGGAG 401

OY 44 AsnPhaIaGluArgIleCysSerGlnValProLysGlySerAsnCysSerAlaSerVal 63
DB 402 ACCATTGTTGGAGAGTCTCGCAAGAAAGTGTGGAAACTTCACTGCGGTGAGAGCGTA 461
OY 64 SerAlaTyrMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSerLeuLysTyr 83
DB 462 ATTGCTTATGTTAAACCGTGTGTGATGAGGCGATTTGTCTGACCGCTTGACAGCATG- 515
OY 84 ProLeuGluAlaLysTyrGlnProLeuThrLeuProAspProTyrGlnLeuGlnAla 103
DB 516 -----AAGTACAAAGCGGTGAGTCTGCAAAATCTTACCAAGTGGAGCGTGC 563
OY 104 PheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSerThrGluLysArgPheTyr 123
DB 564 TTCATGCTTTTACGGAAGATGATCTTAACCTCGCAAGAAATGAGTGAAGCGCTTCGG 623
OY 124 MetArgPheArgArgGlyLysAsnHisSerTyrPheHisAspLeuValPheAsnLeu 143
DB 624 ATCGCTTCGAGG-----AGCAGCCAGCGGCACTACATCACTTGTGTAGCTTGTG 677
OY 144 GluLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsnPhaIaSerArgTyr 163
DB 678 AAGAAAGATGTTGATAGCGACCCCAATCCAATGATGTGAGAACCTTGCATCGCATAC 737
OY 164 LeuTyrMetAlaThrLeuTyrTyrIysThrIysAsnValAspGluPheGlyAlaSer 183
DB 738 TTCTACATGACTACGTTGTACTACAAAGACTTACGACGCTTGACTTACGCGGCTAAG 797
OY 184 PhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrGlyIleLysArgAlaLeu 203
DB 798 TTCTTCAACAAACCTTCTTCAAACTCGCTTGTGTTGTTGATTCAGAAAGCGTTGG 857
OY 204 LysGlnIleIleArgSerAsnLeuProLeuAspIleGlyThrGluHisSerValSerArg 223
DB 856 AAGCGTTGGTTAGAGCAACTTCCCGTGTACCTTGGAAC-CAACCTAGAGCCAC 914
OY 224 LeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIleProAlaLeuPro 243
DB 915 ATCCGCGAAATGATGATAGCGGCTACGCGAGTACATGATGACCCAGCTCGATGAC 974
OY 244 LysPheAlaLysArgPheSerLeuMetValValGlnArgLeuAlaThrValAlaGly 263
DB 975 TCGTTCGCTGAGCGTTTCTCAAGATGGCTACTAGACTCTTGTGTTACCGTCAAG 1034
OY 264 TyrValAspThrProTyrThrLysLysTyrTyrMetLysLeuLysAsnPhaIaAsn 283
DB 1035 TACGTCATTTCCCGCGGTACAAAGAGGTGTACAGAACTTCAAGGAATTCATTGTGAC 1094
OY 284 ArgValPheIleProThrLysLysPheAsnLysGluIleArgGluProSerLys- 302
DB 1095 ---TCTTACAGCACTGCGCAAGTTGATTATGAAGCAAGTCTTCAGCTTAAAGACT 1151
OY 303 AlaLeuLysGluLysValSerThrAspThrLysAspLeuPheGluAsnLysIleGlyGln 322
DB 1152 GCTTACACAAACCTGCTCCCGACAGACACAGGAGCTATCAGGAATGCTCGGTCAA 1211
OY 323 GlyThrValAspPhePheAsnLysGluIleArgAspProSerLysAlaLeuLysGluLys 342
DB 1212 AGCACCAGCAATATGCGCAAC--GGTGTAGCTGATTGTCAAGATGATTAAAGAG- 1265
OY 343 ValSerAsnAspAlaLysAspLeuPheGluAsnLysIle-----Gly 356
DB 1266 -----CCTAAGCCAAACAAATTAATTCGTGAGAGCTGCTCACTACCTTCTAAGCA 1316
OY 357 GlnGlyThrValAspPheIleAsnAsnGluIleArgAsp-----ProSerLysAlaLeu 374
DB 1317 AAGGAGCGCGTTGAGCAAGCTTGTAAAGAGGTTAAATCCGTTGCTCCGATAAG----- 1370
OY 375 IleArgLysValSerThrGlyAlaGluLysPhePheGluAsnLysIleGlyGlnGlyThr 394
DB 1371 ---CAAAAGGCGGACCAACATCCGAGACAGAGCTGTAGAGAAACGTTCCGCTGCG- 1424

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Oy 395 ValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValTyr 414
    ||| ||| ||| |||
Db 1425 ---GATTCGCGGAACGTAATTTGAGTCCCTCGAAGAACAAATCGATCGTGTACT 1481
Oy 415 Thr-----GluAlaAspAsp 419
    ||| ||| ||| |||
Db 1482 ACTCAGAGAGTTAAACAGCAGAGAGTGTATGCCGACGAT 1520

RESULT 4
AAO33064
ID AAO33064 standard; DNA; 1990 BP.
XX
XX AAO33064;
XX
XX 06-MAY-1993 (first entry)
XX
XX Encodes Babesia bovis 60kD immunoreactive merozoite surface epitope.
XX
XX babesiosis; cows; cattle; bos taurus; babesia bovis; babesia bigemina;
XX merozoite; schizont; ss.
XX
XX Babesia bovis.
XX
XX Key Location/Qualifiers
XX CDS 1..1990
XX FT /*tag= a
XX
XX PN US5171685-A.
XX
XX PD 15-DEC-1992.
XX
XX PE 04-APR-1990; 90US-0504461.
XX
XX PR 04-APR-1990; 90US-0504461.
XX
XX PA (USFL ) UNIV FLORIDA.
XX (USDA ) US SEC OF AGRIC.
XX
XX PI Davis WC, Goff WL, Hines SA, Jasmer DP, McElwain TF;
XX Mcgillire TC, Palmergh, Perryman LE, Reduker DW;
XX WPI: 1993-008582/01.
XX DR P-PSDB; AAR30613.
XX
XX PT DNA encoding Babesia bovis protein - is used as probes and for
XX prodn. of polypeptide(s) for use in vaccines and for prodn. of
XX antibodies
XX
XX PS Example 19; Fig 3; 20pp; English.
XX
XX CC This sequence encodes an immunoreactive epitope located on the
XX surface of babesia bovis merozoites. A. B. bovis cDNA expression
XX library was constructed using poly A(+) RNA isolated from B. bovis
XX infected blood cultures. Plaques were immunoscreened using rabbit
XX anti-Bv60 sera, and positive plaques tested for reactivity with
XX monoclonal antibodies that recognised a Bv42 surface exposed epitope
XX as well as an isotype control monoclonal antibody and normal rabbit
XX serum. Lambda rBv60 phagemid DNA was isolated from bacteria, and
XX then restriction digested.
XX
XX SQ Sequence 1990 BP; 628 A; 437 C; 398 G; 527 T; 0 other;

Alignment Scores:
Pred. No.: 1,28e-71 Length: 1990
Score: 826.50 Matches: 178
Percent Similarity: 51.30% Conservative: 79
Best Local Similarity: 35.53% Mismatches: 195
Query Match: 35.04% Indels: 49
Db: 14 Gaps: 6

US-09-807-459-2 (1-458) x AAO33064 (1-1990)
Oy 1 MetaIaProSerAspSerValGlyAspValThrLysThrLeuLeuAlaIaSerGluSer 20

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Db 212 CTCGCTCCAGCTGAAGTGTAGTGAATTATTAACCTCCACATTTGAACAGCTGATCTTG 271
Oy 21 ValAspSerAlaAlaAsnAlaTyrMetIleAsnSerAspMetSerAspTyrLeuSerAla 40
    ||| ||| ||| |||
Db 272 ATGACTCTCCGTGACGACATGACCAACATTACTAGAGATATGAACATGTTTGTGACCAAT 331
Oy 41 ValSerAspAsnPhaIleArgIleCysSerGlnValProLysGluSerAsnCysSer 60
    ||| ||| ||| |||
Db 332 GGTGCGAGCAGATGTGAATGATGTTTGTCTTAATGCTCTGAGACCTCAACTGTCGT 391
Oy 61 AlaSerValSerAlaTyrMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSer 80
    ||| ||| ||| |||
Db 392 GAGGTAGTTAAACAATTATGCTGACCGTTGAAATGATGACGATGCTTACGATTGACAAAT 451
Oy 81 LeuLysTyrProLeuGluAlaLysTyrGlnProLeuThrLeuProAspProTyrGlnLeu 100
    ||| ||| ||| |||
Db 452 GTCAAATATCCGTTGTATCAAGAGTACCAACCTCTATCTCTCCAAACCTTACCGAGTTG 511
Oy 101 GluAlaAlaPheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSerThrGluLys 120
    ||| ||| ||| |||
Db 512 GATGCTGCGCTTCAAGATTTGTTCAAGAGAGTGCATCCAGCCCTGCCAAGAACACGCTRAAA 571
Oy 121 ArgPheTyrMetArgPheArgArgGlyLysAsnHisSerTyrPheHisAspLeuValPhe 140
    ||| ||| ||| |||
Db 572 CGCGAATGCGTTGCGTTTCAACAATGAGCGAACCATGCTGATTACCACTACTTCGTCACAT 631
Oy 141 AsnLeuLeuGluLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsnPhaIle 160
    ||| ||| ||| |||
Db 632 GGTCTGTGTGAACAACATGTTGTGCACAGGAAGAACTCCGATGTGAATATCTTGTTC 691
Oy 161 SerArgTyrLeuTyrMetAlaThrLeuTyrTyrLysThrTyrThrAsnValAspGluPhe 180
    ||| ||| ||| |||
Db 692 AACCAAGTACTTATATGCTGCTACCATGAACTACAAAGCTTATTTGACAGTAACAGATATG 751
Oy 181 GlyAlaSerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrGlyLys 200
    ||| ||| ||| |||
Db 752 AACGCCAAGTTTCAACAACATTCAGCTTCATCAACAAGATTTACGTCGTGATTAG 811
Oy 201 ArgAlaLeuLysGlnIleIleArgSerAsnLeuProLeuAspIleGlyThrGluHisSer 220
    ||| ||| ||| |||
Db 812 CAACAATGTGATGATATCAACAGTGAAGTGTCTCCAAAGATTTT---GAAGAAGAGAGC 868
Oy 221 ValSerArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIlePro 240
    ||| ||| ||| |||
Db 869 ATCGAACGATATCACTCACTACTACTAGCAGCTACGAAAGATTTACTGTTGACCCAGATTCCA 928
Oy 241 AlaLeuProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThr 260
    ||| ||| ||| |||
Db 929 ACCTTTTCCAAAGTTTGACGCTGTTATGCTGACATGCTGAAGAAGTTCGTCGCGTAGC 988
Oy 261 ValAlaGlyTyrValAspThrProTyrTyrLysLysTyrTyrMetLysLeuLysAsnPha 280
    ||| ||| ||| |||
Db 989 TTGACTCGTACGCTTGTAGAGCTCTTGTGTACAAAAGATGATTAAGAATTCACAGACTT 1048
Oy 281 MetValAsnArgValPheIleProThrLysLysPhe----- 292
    ||| ||| ||| |||
Db 1049 TTCTCTAAAAAAGCTTACCCACACTACAAAGAGATTCATCGAGCATCTAACGAAGTTACC 1108
Oy 292 ----- 292

1109 AAAAAGTATCTGAAGCAATGTTGCTGAGCCCACTAAAGTTTATGACGACACTCAC 1168
Oy 293 -----PheAsnLysGluIleArgGluProSerLysAlaLeuLysGlu 306
    ||| ||| ||| |||
Db 1169 GAAAAAACCAAGCTATCTGAAGAAGAAATGTAGCCGAACTCTTAAGACTTTTTCACAG 1228
Oy 307 LysValSerThrAspThrLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAsp 326
    ||| ||| ||| |||
Db 1229 GAGGCTCTCACTCACTCAACCAACTTCTTGATGATGAGAACATTTGCCAACCCACCAAGAG 1288
Oy 327 PhePheAsnLysGluIleArgAspProSerLysAlaLeuLysGluLysValSerAsnAsp 346
    ||| ||| ||| |||

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Db 1289 TTTTTCAGGAAGCTCCCAAGCCACTAAACATTTCTAGACGAAACATCGGTCAACCA 1348
QY 347 AAlaysAspLeuPheGluAsnLysIleGlyInglYThrValAspPheIleAsnGlu 366
Db 1349 ACCAAGGGATTCTTC---AGGAGGCTCTCTCAAGCCACTAAGCCTCTTGGCGAGAAAT 1405
QY 367 ILeArgAspProSerLysAlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPhe 386
Db 1406 ATTGCTCAACCTACTAAAGAAATTTTTCAGAGATGTCCTCAAGTACCAAGAGGTTTATA 1465
QY 387 GluAsnLysIleGlyGlyThrValAspPhe----- 397
Db 1466 ACTGAGAACATTGCTCAACCACTAAGAGATGTCCTGAGGAGAGGTTCTTCATGCTACCATG 1525
QY 398 -----IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValThr 415
Db 1526 AAGTCTTGATGAAGAACATTGCTCAACCTGCGAAGAAATCATATCATGAGTTGTGTACA 1585
QY 416 GluAlaAspAspLeuPheGluAsnLysIleGlyInglYThrValAspPheIleAsnLys 435
Db 1586 GCGCGCAAGAAAT---TTCATTTCCGACAGCCCATGAAAGTACTAAGCAGTTCTTAAACGA 1642
QY 436 GluIleArgAspProSerLysAlaLeuIleArg---LysValSerThrGluAlaAspAsn 454
Db 1643 ACTGTGGCCAACTTCAAGAGAAATTCCTGAACGAGGCTTTAGAAACATTAAGACGCA 1702
QY 455 Leu 455
Db 1703 TTA 1705

RESULT 5
AAT18995
ID AAT18995 standard; cDNA; 1990 BP.
XX
AC AAT18995;
XX
DT 15-OCT-1996 (first entry)
XX
DE Babesia merozoitae surface protein cDNA clone Bv60.
XX
KM Babesia; merozoitae protein; vaccine; probe; diagnosis; ss.
XX
OS Babesia bovis.
XX
FH Key Location/Qualifiers
FT CDS 122..1819
FT /tag= a
XX
FN US518916-A.
XX
PD 21-MAY-1996.
XX
PF 04-APR-1989; 89US-0333155.
XX
PR 04-APR-1990; 90US-0504461.
PR 04-APR-1989; 89US-0333155.
PR 14-DEC-1992; 92US-0989616.
PR 21-NOV-1994; 94US-0342480.
XX
PA (USDA ) US SEC OF AGRIC.
XX
PI Goff WL, Jasmer DP, McElwain TF, McGuire TC, Reduker DW;
PI Stillier D;
XX
DR WPI: 1996-259067/26.
DR P-PSDB: AAR97981.
XX
PT New fragment of Babesia bovis genomic DNA - useful as a probe for
PT detecting Babesia infection
XX
PS Example 18; Fig 3; 19pp; English.
XX
CC A cDNA clone (AAT18995) codes for Bv60, a 60 kDa immunoreactive

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CC protein (AAR97981) located on the surface of Babesia bovis merozoites.
CC It was isolated from a blood-stage B. bovis cDNA library in lambda
CC ZAP1 by subcloning into Bluescript SK(-) and immunoscreening using
CC monospecific anti-Bv60 antiserum. Bv60, Bv44 and Bv42 (see also
CC AAT18993 and AAT18994) DNA sequences can be used to make recombinant
CC proteins useful as vaccines for the prophylaxis of bovine babesiosis.
CC They can also be used as diagnostic probes.
XX
SQ Sequence 1990 BP; 628 A; 437 C; 398 G; 527 T; 0 other.

Alignment Scores:
Pred. No.: 1,28e-71 Length: 1990
Score: 826.50 Matches: 178
Percent Similarity: 51.30% Conservative: 79
Best Local Similarity: 35.53% Mismatches: 195
Query Match: 35.04% Indels: 49
DB: 17 Gaps: 6

US-09-807-459-2 (1-458) x AAT18995 (1-1990)
QY 1 MetaIAspProSerAspSerValGlyAspValThrLysThrLeuLeuAlaSerGluSer 20
Db 212 CTCGCTCCAGCTGAGGTGAGTGTAGCTATTTACCTCCATTTGGAAACAGCTGATCTTG 271
QY 21 ValAspSerAlaAlaAsnAlaTyrMetIleAsnSerAspMetSerAspPyrLeuSerAla 40
Db 272 ATGACTCTCCGTGACCACTGACCAACATTTACTAGATATGAACATGTTTGGCAAT 331
QY 41 ValSerAspAsnPheAlaGluArgIleCysSerGlnValProlyGlySerAsnGlySer 60
Db 332 GGTCTGTGACAGATTGTAAATGATGTTGCTTAATGCTCTCCAGAGACTCCACAGCTGCT 391
QY 61 AlaSerValSerAlaTyrMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSer 80
Db 392 GAGGTAGTTAACAAATTATGCTGACCGCTGTGAATATGATGATGCTTACGATTGACAT 451
QY 81 LeuLysTyrProLeuGluAlaLysTyrGlnProLeuThrLeuProAspProTyrGlnLeu 100
Db 452 GCGAAATATCCGTTGTATCAAGAGTACCAACCTCTATCTCTCCAAACCTTACCACTTG 511
QY 101 GluAlaAlaPheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSerThrGluLys 120
Db 512 GATGCTGCTGATGATGTTGTTCAAGAGAGTGCATGAAACCTGCCAAGACAGCTTAAA 571
QY 121 ArgPheTyrMetLeuArgPheArgGlyLysAsnHisSerTyrPheHisAspLeuValPhe 140
Db 572 CGCGAATGTTGCGTTTCAGAAATGAGACGAACATGATGATTACCTACTGCTGACT 631
QY 141 AsnLeuLeuGluLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsnPheAla 160
Db 632 GGTCTGTGACCAACAAATGTTGTCGACGAGCAAGCACTACCGATGTAATCTTGTTC 691
QY 161 SerArgTyrLeuTyrMetAlaThrLeuTyrTyrLysThrTyrThrAsnValAspLuphe 180
Db 692 AACCAAGTACTATATGCTATGACTACATGAACTACAGCTATTTGACAGTAACGATATG 751
QY 181 GlyAlaSerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrDpLysLys 200
Db 752 AACGCAAGTCTTTCACAGATTCAGCTTCACTACAAAGATATTCAGTCGATTTAGG 811
QY 201 ArgAlaLeuLysGlnIleIleArgSerAsnLeuProLeuAspIleGlyThrGluHisSer 220
Db 812 CAACATATGATGATATCATGAGTGTCTCTCAAGATTTT---CAAGAAAGAGAC 868
QY 221 ValSerArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIlePro 240
Db 869 ATCGAAGCTATCACTCAACCTTACTAGACAGTACGAAGTATCACTGTGACCCAGATTCCA 928
QY 241 AlaLeuProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThr 260
Db 929 ACTCTTCCAAAGTTTGACGCTGATATGCTGACATGATGGAAGAGCTTCTGCTGATGC 988
QY 261 ValAlaGlyTyrValAspThrProTyrTyrLysTyrTyrMetLysLeuLysAsnPhe 280

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Db 989 TTGACCTGCTAGCTGAGCTCTTGGTACAAAGATGATTAAGAAATTGAGACTTT 1048
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 281 MetValAsnArgValPheIleProThrLysLysPhe----- 292
      ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1049 TTCCTAAAGCGTTACCAACTTACAAAGAGTTCTATCGAGATTAACGAGTTACC 1108
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 292 ----- 292
Db 1109 AAAAATCTCTGAAGCCAAATCTTCTGAGCCACTAAAGTTATATGACGACACTC 1168
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 293 -----PheAsnLysGluIleArgGluProSerLysAlaLeuLysGlu 306
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1169 GAAAAACCAAGGCTATCTGAAAGATGATGAGCAAGCAACTTAAAGACTTTTTCAG 1228
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 307 LysValSerThrAspThrLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAsp 326
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1229 GAGGCTCCTCAAGTCAACCACTTCTTCATGAGAACATTGGCCACCCCAAGAG 1288
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 327 PhePheAsnLysGluIleArgAspProSerLysAlaLeuLysGluLysValSerAsnAsp 346
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1289 TTTTTCAGGGAAGCTCCCAACCCACTAAACATTCTTAGACGAAACATCGCTACCA 1348
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 347 AlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnAsnGlu 366
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1349 ACCAAGGAGTCTTC--AGGAGAGCTCCTCAAGCCACTAAGCAGCTCTCTAGCGCAGAT 1405
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 367 IleArgAspProSerLysAlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPhe 386
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1406 ATTGCTCAACCTACTAAAGATTTTTCAAAGATGTCCCTCAAGTCAACCAAGAGTTATA 1465
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 387 GluAsnLysIleGlyGlnGlyThrValAspPhe----- 397
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1466 ACTGGAACATTGCTCAACCACTAAGAGGTTCCGAGAGGAGGTTCTCATCCTACCATG 1525
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 398 -----IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValThr 415
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1526 AAAGTCTGAATGAAGAAATTCCTCACTGCGCAAGAAATCATGATGATTTGGTACA 1585
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 416 GluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnLys 435
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1586 GCGGCCAAGAAAT--TTCATTTCCCGACGCCATGAGAGTACTAACAGTTCTTAAACGAA 1642
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 436 GluIleArgAspProSerLysAlaLeuIleArg--LysValSerThrGluLysAsn 454
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1643 ACTGTGGCCAACTTACAAAGAAATTCCTGAACGAGCTTTAGAACTTACTAAAGACGA 1702
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 455 Leu 455
      |||
Db 1703 TTA 1705

RESULT 6
AA047076
ID AA047076 standard; DNA; 1371 BP.
XX
AC AA047076;
XX
DE 13-JAN-1994 (first entry)
XX
DE B. canis 21B4/rhoptry antigen gene 2 DNA.
XX
KW Polymerase chain reaction; PCR; amplify; primer; detection;
KW babesiosis; parasite; Babesia bovis; 21B4/rhoptry; antigen; gene;
KW repeat region; immune response; vaccine; ss.
OS Babesia canis.
XX
PN WO9314204-A.
XX
PD 22-JUL-1993.
XX
PF 15-JAN-1993; 93WO-A000012.
XX

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PR 15-JAN-1992; 92AU-0000399.
XX
PA (CSTR ) COMMONWEALTH SCI & IND RES ORG.
XX
PI Dalrymple BP, Peters JM;
XX
DR WPI; 1993-243219/30.
DR P-PSDB; AAR39902.
XX
PR Detecting closely linked gene copies which encode protective
PR antigen against babesiosis - by screening babesial genomic DNA
PR library with oligo-nucleotide probe based partial sequencing of
PR protective antigen and identifying positive clones
XX
PS Claim 24; Fig 7; 55pp; English.
XX
CC This sequence represents the Babesia canis 21B4/rhoptry antigen gene
CC 2. This sequence was determined from restriction fragments from the
CC clone B. canis lambda GBM-11 #9. B. canis was found to contain two
CC genes which are related to the B. bovis 21B4 gene. Gene 1 and gene 2
CC are very similar but gene 2 appears to contain a large number of
CC repeats. Babesia antigen genes can be used in the production of a
CC combined vaccine which will stimulate a greater immune response and
CC afford broader immunity than a single antigen vaccine. See also
CC AA047068-74.
XX
SQ Sequence 1371 BP; 418 A; 312 C; 325 G; 318 T; 0 other;
Alignment Scores:
Pred. No.: 2,556-66 Length: 1371
Score: 770.50 Matches: 161
Percent Similarity: 55.65% Conservative: 95
Best Local Similarity: 35.00% Mismatches: 161
Query Match: 32.66% Indels: 43
DB: 14 Gaps: 9
US-09-807-459-2 (1-458) x AA047076 (1-1371)
Qy 1 MetAlaProSerAspSerValIleGlyAspValThrLysThrLeuLeuAlaIleSerGluSer 20
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 91 CTTTCTAAATCAGATGAGAGCCGAAAGACCTTGTCTTCACTCTTCACTTACGTCGACCATG 150
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 21 ValAspSerAlaAlaAsnAlaThrMetIleAsnSerAspMetSerAspThrLeuSerAla 40
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 151 ACGAGAGCGGCTTTAGAACGCTACAGAAATGATGCTGATGCAACATTTTCAACCGT 210
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 41 ValSerAspAsnPheAlaGluArgIleCysSerGlnValProLysGlySerAsnCysSer 60
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 211 CGCAGGGAAGAGAGAGAGAGAGCTGCTGTGGAACATCGCAGAGAGAGACTGAATGTGAG 270
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 61 AlaSerValSerAlaThrMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSer 80
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 271 AAGAGCGTAGCGTAGTATGTTGAAAGCTCGTCAGGTACGACTGTTGACATGGAAC 330
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 81 LeuLysTyProLeuGluAlaLysTyArgInProLeuThrLeuProAspProTyArgInLeu 100
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 331 CAGAACTACCTCAGAGAGAAAGATACAGCCGCTACCTCCCAACCTTATCATATG 390
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 101 GluAlaAlaPheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSerThrGluLys 120
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 391 GAGGCCGCGTTCTATGCTTTCAGAAACAGTGAATCAAACTTAAACCAACCAACGAA 450
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 121 ArgPheThrMetArgPheArgArgGlyLysAsnHisSerTyPheHisAspLeuValPhe 140
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 451 GCTTCTGGATCGTTTCTGTCACGCGGAGCTTATGCGCCCTATACCAATTCCTCGGTG 510
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 141 AsnLeuLeuGluLysAsnValThrArgAspAlaAspAlaThrAspIleGluLysPheAla 160
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 511 AATATTCTATATAAAACCTCAGCGATACATGCGATGATGATACCTCAGAGTTTCGTA 570
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 161 SerArgTyLeuTyMetAlaThrLeuTyTyLysThyTyThrAsnValAspGluPhe 180
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 571 CGCAATATGCTTACATGCGCACCATGTATATACAGACATACACCGCTTGTGATGTGTA 630
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::

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OY 121 ArgPheTrpMetArgPheArgArgGlyLysAsnHisSerTyrPheHisAspLeuValPhe 140
DB 451 GGTCCATGGATGGTACAGAGAGGAGGATGAACATGGTACATTCATCCATCA 510
OY 141 AsnLeuLeuGlnLysAsnValThrArgAspAlaAspAlaThrAspIleGlnAsnPheAla 160
DB 511 AGTTTGCTCGGCAGAGATGGTTCCTAAGATGGTGTACTGACTTGACTCCCGTC 570
OY 161 SerArgTyrLeuTyrMetLalaThrLeuTyrTyrLysThrTyrThrAsnValAspGluPhe 180
DB 571 AACAGCTTTGGTACATGGCCACACCTACTACAAAACCTACTAATGTCAAGAGATTG 630
OY 181 GlyAlaSerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyIlePglyIleLys 200
DB 631 GGTGCTCGCTTTTCAACACTTCTCTTTACAAATATATATCGGTATGATTTAA 690
OY 201 ArgAlaLeuLysGlnIleLeuArgSerAsnLeuProLeuAspIleGlyThrGlnHisSer 220
DB 691 AGGGCATTAAGAGCATGCTCGCTCCATGTTCCGGAAGACATGGGA---GAGCAGAT 747
OY 221 ValSerArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIlePro 240
DB 748 ATTGAACGTATGATCATTTGTCGGAAGGATACAGACTACATGTTGACACAGCTGCCA 807
OY 241 AlaLeuProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThr 260
DB 808 ACCCTTTCAAGATTGTCGCAACGTTACTGACATGTTATGAACTTCTGTTGACAGC 867
OY 261 ValAlaGlyTyrValAspThrProTyrTyrLysLysTrpTyrMetLysLeuLysAsnPhe 280
DB 868 CTCGCCGGTTATGTCAAGGCTCATGTTCAAGAGATGATGATTAATGATGTCCTTG 927
OY 281 MetValAsnArg-----ValPheIle 287
DB 928 TTAACGTGTGAAGCTTACCAACCTGTATGAGATATACATTACTTAACCTATTTCGTT 987
OY 288 ProThrLysLysPhePheAsnLysGlnIleArgGluProSerLys---AlaLeuLysGlu 306
DB 988 GATACCTCCAGGAATTAATCAATAAGATGACATTAAACCTGCTGATGCTGAGAGAA 1047
OY 307 LysValSerThrAspThrLysAspLeuPheGlnAsnLysIleGlyGlnGlyThrValAsp 326
DB 1048 AATATCGTTAACCCCTGATGTTATCTCCGACGCAACAAATATTTCTAGTGCACA 1107
OY 327 PhePheAsn-----LysGlnIleArgAspProSerLysAlaLeu 339
DB 1108 AACTACAAATGACGGCATATAAATAGATCCCTCTTTATGAACTTAAGAGCGGCAT 1167
OY 340 LysGlnLysValSerAsnAspAlaLysAspLeuPheGlnAsnLysIleGlyGlnGlyThr 359
DB 1168 ATCCGAATTGCTGCGAATACCGCTAGCATTAATAGATGATTAAGTAATAA----- 1221
OY 360 ValAspPheIleAsnAsnGlnIleArgAspProSerLysAlaLeuIleArgLysValSer 379
DB 1222 -----GCGAAAGAAATTAAGTGCCTGCCAAG 1248
OY 380 ThrGlyAlaGluAspLeuPheGlnAsnLysIleGlyGlnGlyThrValAspPheIleAsn 399
DB 1249 GACCGCGCAACAGCATATATGACAGATACAGTAAACCTGCTGTAGTGTATTAACGAC 1308
OY 400 -----AsnGlnIleArgAspProSerLysAlaLeuIleArgLysValTyr----- 414
DB 1309 GTTGTAAAGAAATGATCTCTTGTATGACAGTAAC-----ATCAGAAATATATTAACGTGC 1362
OY 415 ---ThrGlnAlaAspAspLeuPheGlnAsnLysIleGlyGlnGlyThrValAspPheIle 433
DB 1363 AGCTTCAAGATGACAAATATATGAACAAGAGACTGAGAGAGAGAAAGTTGAGGAAGTT 1422
OY 434 AsnLysGlnIleArgAspProSerLysAla 443
DB 1423 AAACCTGACCTGAAGCAAAAAGATGTGCT 1452

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RESULT 8
AAQ26065
ID AAQ26065 standard; DNA; 282 BP.
XX
AC AAQ26065;
XX
DE 09-DEC-1992 (first entry)
XX
DE 21B4 gene clone pT#13, EcoRI insert.
XX
KM Beta-galactosidase; B. bovis; Bb; T21B4; ss.
XX
OS Babesia bovis.
XX
PN EP492525-A.
XX
PD 01-JUL-1992.
XX
PF 20-DEC-1991; 91BP-0121990.
XX
PR 21-DEC-1990; 90AU-0004051.
XX
PA (CSIR ) COMMONWEALTH SCI & IND RES ORG.
XX
PI Casu RE, Commins MA;
XX
DR WPI, 1992-218727/27.
XX
DR P-PSDB; AAR25187.
XX
PT Monoclonal antibody to Babesia bovis parasite - used to isolate
PT antigens for use in vaccines for treating Babesiosis and
PT providing immunity in cattle
XX
PS Claim 13; Fig 9; 24pp; English.
XX
CC The sequences given in AAQ26062 and AAQ26065-7 are portions of the 21B4
CC gene which were isolated from a B. bovis (Bb) cDNA lambda gt11
CC library and cloned into pGEM7zf(+). The resulting plasmids were
CC transformed into E. coli strain JM83. The inserts were in frame,
CC when translated, with the vector beta-galactosidase gene. The fusion
CC proteins produced by translation of these vectors were recognised by
CC the monoclonal antibody of the invention, T21B4. These fusion
CC antigens could be used in vaccines for the treatment of babesiosis
CC and to provide immunity in relation to Bb infection in cattle against
CC different strains of Babesia by heterologous and homologous challenge.
XX
SQ Sequence 282 BP; 80 A; 60 C; 63 G; 79 T; 0 other;

Alignment Scores:
Pred. No.: 3,18e-07 Length: 282
Score: 161.00 Matches: 32
Percent Similarity: 51.65% Conservative: 15
Best Local Similarity: 35.16% Mismatches: 44
Query Match: 6.82% Indels: 0
Gaps: 0

US-09-807-459-2 (1-458) x AAQ26065 (1-282)
OY 2 AlaProSerAspSerValGlyAspValThrLysThrLeuLeuAlaIleSerGluSerVal 21
DB 10 GCTCCAGCTGAAAGTGAAGTGAATTAATCTCCACATTTGAAACAGCTGATTAATTGATG 69
OY 22 AspSerAlaIleAsnAlaTyrMetIleAsnSerAspMetSerAspTyrIleSerAlaVal 41
DB 70 ACTCTCGTGACCAACATGCAACAACTTACTAAGAGATGAAACAGCTTTGACCAATGCT 129
OY 42 SerAspAsnPheAlaGluArgIleCysSerGlnValaProLysGlySerAsnCysSerAla 61
DB 130 CGTGACAGATGTGAATGATGTTTGGCTTAATGCTCTGAGGACCTCCAACTGTCGTAG 189
OY 62 SerValSerIleArgLysSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSerLeu 81
DB 190 GTAGTTAAACAACATATGCTGACCGCTTGAAGATGATGACGATTCACGATTCACAAATGTC 249

```

```

0Y      82 LysTyProLeuGluAlaLysTyGInProLeu 92
      ::::::::::::::::::::
Db      250 AGATATCGGTGATCCAGAGTACCAACCTCTA 282

RESULT 9
AAV21209
ID      AAV21209 standard; DNA; 1664976 BP.
XX
XX      AAV21209;
AC
XX      10-NOV-1998 (first entry)
XX
XX      Methanococcus jannaschii circular chromosome.
DE
XX      Methanococcus jannaschii; methanogenic archaeon; circular chromosome;
XX      genome; autotrophic; extrachromosomal element; identification; ds.
XX
XX      Methanococcus jannaschii.
OS
XX      PN
XX      WO9807830-A2.
XX
XX      26-FEB-1998.
PD
XX      22-AUG-1997; 97WO-US14900.
XX
XX      22-AUG-1996; 96US-0024428.
XX
XX      (GENO-) INST GENOMIC RES.
XX      (UNIT) UNIV ILLINOIS FOUND.
XX      (UYXO ) UNIV JOHNS HOPKINS SCHOOL MEDICINE.
XX
XX      Bult CJ, Smith HO, Venter JC, White OR, Woese CR;
XX      WPI: 1998-169145/15.
XX
XX      Complete genome sequence of methano-genic archaeon, Methanococcus
XX      jannaschii - useful in identification of M. jannaschii genome
XX      fragment
XX
XX      Claim 13; Page 152-585; 614pp; English.
XX
XX      The present sequence represents the complete 1.66-megabase pair genome
XX      sequence of the Methanococcus jannaschii circular chromosome. The
XX      present invention describes M. jannaschii open reading frames from the
XX      genome sequence. The invention also describes a computer based system
XX      for identifying fragments of the M. jannaschii genome that are
XX      homologous to target nucleotide sequences, comprising: (a) data storage
XX      means comprising the nucleotide sequence of the 1664976, 58407 or 16550
XX      bp sequence (see AAV21209, AAV21210 and AAV21211), or a nucleotide
XX      sequence at least 99.9% identical to it; (b) search means for comparing a
XX      target sequence to the nucleotide sequence of the data storage means to
XX      identify a homologous sequence, and (c) retrieval means for obtaining
XX      the homologous sequence. The method, which is based on whole genome
XX      random sequencing of an autotrophic archaeon M. jannaschii, the genome
XX      of which consists of 3 physically distinct elements, a large circular
XX      chromosome (the 1664976 bp sequence given in AAV21209), a large circular
XX      extra-chromosomal element (the 58407 bp sequence given in AAV21210), and
XX      a small circular extra-chromosomal element (the 16550 bp sequence given
XX      in AAV21211), can be used in the identification of M. jannaschii genome
XX      fragment.
XX
XX      Sequence 1664976 BP; 568133 A; 264649 C; 258701 G; 573392 T; 101 other;
XX
Alignment Scores:
Pred. No.: 0.823 Length: 1664976
Score: 147.00 Matches: 88
Percent Similarity: 38.79% Conservative: 59
Best Local Similarity: 23.22% Mismatches: 150
Query Match: 6.23% Indels: 82
DB: 19 Gaps: 17

US-09-807-459-2 (1-458) x AAV21209 (1-1664976)

```

Oy 440 roSerLyAlaLeuIleArgLyValSerThrGluAlaAspAsnIleuGlu 457
Db 1419240 TTGGGGGAGAAATTGTTAAATAATACATTAAGTCTTAGAG--TTGTTAGAG 1419289
RESULT 10
ID AAQ26066 standard; DNA: 170 BP.
XX
AC AAQ26066;
XX
DT 09-DEC-1992 (first entry)
XX
DE 21B4 gene clone pT#13, EcoRI insert (2).
XX
KM Beta-galactosidase; B. bovis; Bb; T21B4; ss.
XX
OS Babesia bovis.
XX
PN EP492525-A.
XX
PD 01-JUL-1992.
XX
PF 20-DEC-1991; 91EP-0121990.
XX
PR 21-DEC-1990; 90AU-0004051.
XX
PA (CSIR) COMMONWEALTH SCI & IND RES ORG.
XX
PI Casu RE, Commings MA;
XX
DR WPI: 1992-218727/27.
XX
P-PSDB: AAR25188.
XX
PT Monoclonal antibody to Babesia bovis parasite - used to isolate
PT antigens for use in vaccines for treating Babesiosis and
PT providing immunity in cattle
XX
PS Claim 15; Fig 10; 24pp; English.
XX
CC The sequences given in AAQ26062 and AAQ26065-7 are portions of the 21B4
CC gene which were isolated from a B. bovis (Bb) CDNA lambda dbt1
CC library and cloned into pGEM7zf(+). The resulting plasmids were
CC transformed into E. coli strain JM83. The inserts were in frame,
CC when translated, with the vector beta-galactosidase gene. The fusion
CC proteins produced by translation of these vectors were recognised by
CC the monoclonal antibody of the invention, T21B4. These fusion
CC antigens could be used in vaccines for the treatment of babesiosis
CC and to provide immunity in relation to Bb infection in cattle against
CC different strains of Babesia by heterologous and homologous challenge.
XX
SQ Sequence 170 BP; 56 A; 28 C; 31 G; 51 T; 4 other;
XX
Alignment Scores:
Pred. No.: 2.98e-05 Length: 170
Score: 138.00 Matches: 27
Percent Similarity: 66.67% Conservative: 9
Best Local Similarity: 50.00% Mismatches: 18
Query Match: 5.85 Indels: 0
Gaps: 0
US-09-807-459-2 (1-458) x AAQ26066 (1-170)
Oy 161 SerArgTyrlLeuTyrlMetAlaThrLeuTyrlTyrlThrAsnValAspGluPhe 180
Db 1 AACAGGTAAGTCTTAATAGCTACATGACGTAAGACTTATTGACAGTAACAGTAG 60
Oy 181 GYAASerPhePheAsnIleuSerPheThrThrGlyLeuPheGlyTrpGlyIleLys 200
Db 61 AACGCCAANTTCTTCAACAGATTCAGCTTACTACAAAGATTTTCAGTNNTCGTATTAG 120
Oy 201 ArgAlaLeuLysGlnIleIleArgSerAsnLeuProLeuAsp 214
Db 121 CAATCATTTGAGTATATCATCATCGTGAATGTCTCTGAAGAT 162

RESULT 11
ID AAX20254/c standard; DNA: 26776 BP.
XX
AC AAX20254;
XX
DT 04-MAY-1999 (first entry)
XX
DE Borrelia burgdorferi polynucleotide sequence #7.
XX
KM Borrelia burgdorferi; spirochete; bacterium; pathogen; Lyme disease;
KM epidemic relapsing fever; endemic relapsing fever; Lyme borreliosis;
KM infection; diagnosis; characterisation; detection; ds.
XX
OS Borrelia burgdorferi.
XX
PN W09858943-A1.
XX
PD 30-DEC-1998.
XX
PF 18-JUN-1998; 98WO-US12764.
XX
PR 03-SEP-1997; 97US-0057483.
PR 20-JUN-1997; 97US-0050359.
PR 22-JUL-1997; 97US-0053344.
PR 22-JUL-1997; 97US-0053377.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PA (MEDI-) MEDIMMUNE INC.
XX
PI Clayton R, Dougherty BA, Fraser C, Lathigra R, Smith HO;
XX
PI White OR;
XX
DR WPI: 1999-081217/07.
XX
PT New isolated Borrelia burgdorferi nucleic acids - used to develop
PT products for the detection, diagnosis, characterisation, prevention
PT and therapy of infections, particularly Lyme disease
XX
PS Claim 1; Page 867-882; 1128pp; English.
XX
XX AAX20248 to AAX20402 represent polynucleotide sequences isolated from
XX Borrelia burgdorferi (Bb). Products derived from Bb can be used for
XX the detection, diagnosis, characterisation, prevention and therapy of
XX Bb infections, e.g. Lyme disease. They can also be used for the
XX production of biosynthetic products, e.g. enzymes. Borrelia belongs
XX to a family of motile, spiral-shaped bacteria called Spirochetes.
XX Spirochetes are pathogenic in humans and Borrelia causes epidemic and
XX endemic relapsing fever, and Lyme borreliosis, more commonly known as
XX Lyme disease.
XX
SQ Sequence 26776 BP; 8604 A; 4331 C; 4257 G; 9580 T; 4 other;
XX
Alignment Scores:
Pred. No.: 0.0633 Length: 26776
Score: 134.00 Matches: 61
Percent Similarity: 40.96% Conservative: 75
Best Local Similarity: 18.37% Mismatches: 124
Query Match: 5.68 Indels: 72
Gaps: 13
US-09-807-459-2 (1-458) x AAX20254 (1-26776)
Oy 167 AlaThrLeuTyrlTyrlLeuTyrlThrAsnValAspGluPheGlyAlaSerPhePheAsn 186
Db 6709 GCTACAAATTAAGTCTTAACATC-----AAATTGGGTGAG 6677
Oy 187 LysLeuSerPheThrThrGlyLeuPheGlyTrpGlyIleLysArgAlaLeuLysGlnIle 206
Db 6676 CTGTAGCATATCTTCCCGCAAGTGGTGAAGGCAATGATAATGTCTTCAAGGCCAATT 6617
Oy 207 IleArgSerAsnLeuProLeuAspIleGlyThrGlnHisSerValSerArgLeuGlnHis 226

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Db 6616 ACCAGCCCTTTGTTAAAGCTTACGAGGAATCAATACGTAACGGCAACCTAAAGAA 6557
Oy 227 ILehrserSerThrLysAspThrMetAspThrGlnIleProAlaLeuProLysPheAla 246
Db 6556 CTTGCAACA-----TATGTTCTTAGTGATTT-----TTAGATCGTTTAAAG 6515
Oy 247 LysArgPheSerLeuMetVal-----ValGlnArgLeuLeuAlaThrValAlaGlyTyr 264
Db 6514 GGTGCGTTTGAGCCATTATCAATGGGTTCAAAAGTTGATGATGATATCC----- 6461
Oy 265 ValAspThrProTrpTyrLysLysTrpMetLysLeuLysAsnPheMetVal----- 282
Db 6460 -----AAAGCGTATACAAATTTGAAATTCGTGTTACTTTTGG 6422
Oy 283 -----AsnArgValPhe 286
Db 6421 AAAAAGCCGAAAGAAAGAAAGACATTCTGCTCCAGACCCGAAAGGATTAAGTTT 6362
Oy 287 ILeProThrLysLysPhe---PheAsnLysGlnIleArgLysProSerLysAlaLeuLys 305
Db 6361 GATCCAATGCAAAACCAATATTAACAGAAATGGCAGAAACCTATCAAAACCTGCAA 6302
Oy 306 GlnLysValSerThrAspThrLysAspLeuPheGlnAsnLysIleGlyGlnGly----- 323
Db 6301 GACGAATATTTAATAGCGCAGAGGACATCTAT--AATAAGACCGGGAAGACAGAGAA 6245
Oy 324 -----ThrValAspPhePheAsnLysGlnIleArgAspPro 335
Db 6244 CAAGCCTTAAGATCTTGAAAAAACCATTAATGAAAAAACCAAAAGTTTAAGATGAA 6185
Oy 336 SerLysAlaLeuLysGlnLysValSerAsnAspAlaLysAspLeuPhe----- 351
Db 6184 TACTCCAAATATTTGATTCAGTTAACTGACGAGAAACAAATTTAGTCGAGTTGAA 6125
Oy 352 -----GlnAsnLysIleGlyGlnGlyThrValAspPheIleAsnGlnIleArgAsp 369
Db 6124 AATCACTAATTAATGATTCATTAACCTCAATTAATGATTTTGGAATGAG----- 6077
Oy 370 ProSerLysAlaLeuIleArgLysValSerThrGlnLysAlaGlnAspLeuPheGlnAsnLys 389
Db 6076 ---TACCAAAATTTACTTAAAGAAAGAAAGAGTGTGAAGGAAATATATCAAAACTCG 6020
Oy 390 IleGlyGlnGlyThrValAspPhe-----IleAsnAsnGlnIleArgAspProSer 406
Db 6019 CCCCATACAGATCAAGTACGCGCTTTACAGAAACCTTAATGATGAAATCAATGAAAGAAC 5960
Oy 407 LysAlaLeuIleArgLysValTyrThrGlnLysAspAspLeuPheGlnAsnLysIleGly 426
Db 5959 AAAGCGTTTCGTAAGAAATATGAAAAAGTTTCGAAACTCG-----AACGAGCTTAAC 5906
Oy 427 GlnGlyThrValAspPheIleAsnLysGlnIleArgAspProSerLysAlaLeuIleArg 446
Db 5905 AGGCATAGTTAGTCGACCTTGAAGAACAGGTTATATGAAATGAAAAACCGCTTTGAT 5846
Oy 447 LysValSerThrGlnLysAspAsnLeuLeuGlnLys 458
Db 5845 CGATCTTTTGTGCGAGCTCAAAAAAGCTCTGCAAAA 5810

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RESULT 12
AAx20250/c
ID AAx20250 standard; DNA: 111309 BP.

AAx20250;
AC
XX
DT 04-MAY-1999 (first entry)

DE Borrelia burgdorferi polynucleotide sequence #3.

XX
KW Borrelia burgdorferi; spirochete; bacterium; pathogen; Lyme disease;
epidemic relapsing fever; endemic relapsing fever; Lyme borreliosis;
infection; diagnosis; characterisation; detection; ds.

```

OS Borrelia burgdorferi.
PN
XX MO9858943-A1.
XX
XX 30-DEC-1998.
XX
XX 18-JUN-1998; 98WO-US12764.
XX
XX 03-SEP-1997; 97US-0057483.
XX 20-JUN-1997; 97US-0050359.
XX 22-JUL-1997; 97US-0053344.
XX 22-JUL-1997; 97US-0053377.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX (MEDI-) MEDIMUNE INC.
XX
XX Clayton R, Dougherty BA, Fraser C, Lethigra R, Smith HO:
XX white OR;
XX
XX WPI; 1999-081217/07.
XX
XX New isolated Borrelia burgdorferi nucleic acids - used to develop
XX products for the detection, diagnosis, characterisation, prevention
XX and therapy of infections, particularly Lyme disease
XX
XX Claim 1; Page 738-800; 1128pp; English.
XX
XX AAx20248 to AAx20402 represent polynucleotide sequences isolated from
XX CC Borrelia burgdorferi (Bb). Products derived from Bb can be used for
XX the detection, diagnosis, characterisation, prevention and therapy of
XX CC Bb infections, e.g. Lyme disease. They can also be used for the
XX CC production of biosynthetic products, e.g. enzymes. Borrelia belongs
XX CC to a family of motile, spiral-shaped bacteria called Spirochetes.
XX CC Spirochetes are pathogenic in humans and Borrelia causes epidemic and
XX CC endemic relapsing fever, and Lyme borreliosis, more commonly known as
XX CC Lyme disease.
XX
XX SQ Sequence 111309 BP; 35956 A; 13151 C; 19075 G; 43117 T; 10 other:
XX
XX
XX Alignment Scores:
XX Pred. No.: 5.75 Length: 111309
XX Score: 122.50 Matches: 93
XX Percent Similarity: 37.47% Conservative: 55
XX Best Local Similarity: 23.54% Mismatches: 144
XX Query Match: 5.19% Gaps: 103
XX DB: 20
XX
XX US-09-807-459-2 (1-458) x AAx20250 (1-111309)
XX
XX Oy 113 AsnProAlaAsnSerThrGlnLysArgPheTrpMetArgPheArgGlyLysAsnHis 132
XX |||||
XX Db 47193 AACCCATTAACCAAGAAATTAACACATTT----- 47164
XX
XX Oy 133 SerYrPheHisAspLeuValPheAsnLeuLeuGlnLysAsnValThrArgAspAlaAsp 152
XX |||||
XX Db 47163 AAACCTTTTAAATGATGATTTTATATCTTTAAAGAAATTTTAAACAGGAA----- 47110
XX
XX Oy 153 AlaThrAspIleGlnAsnPheAlaSerArgTyrLeuTyrMetAlaThrLeuTyrTyrLys 172
XX |||||
XX Db 47109 -----GCCCTAGAAATATTTTAAAGAAAGCTTTTCGATTAATAAAA 47071
XX
XX Oy 173 ThrTyrThrAsnValAspGluPhe---GlyAlaSerPhePheAsnLysLeuSerPheThr 191
XX |||||
XX Db 47070 ACAGATGAAGATGCMAATTAATATCTTGCTGACATTTTAAAGAAATGATATAAA 47011
XX
XX Oy 192 ThrGlyLeuPheGlyTyrPglyTlleLysArgAlaLeuLysGlnIle----- 206
XX |||||
XX Db 47010 GAACCTCTATAC-----TATCTTAAAAAGTCCCAATTTTAATAAA 46969
XX
XX Oy 207 -----IleArgSerAsnLeuProLeuAspIleGlyThrGlnHisSerValSer 222
XX |||||
XX Db 46968 AATAATAAGATGTCGTAACCAACCTGGAAATTAACCTTATTAATCTAGAAAGCTACGA 46909

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QY	263	GLYThrValAspThrProTrrPTyLysLysStrPtyrMetLysLeuLysAsnPhmeVal	282
Db	636234	TTGGCAATTCGCTAAATCTTTATACAAAATAATCAACAGATCATCTACGACTAGAGATTGGC	636175
QY	263	GLYThrValAspThrProTrrPTyLysLysStrPtyrMetLysLeuLysAsnPhmeVal	282
Db	636174	AAACAAATVA	636145
QY	263	AsnArgValPheIleProThrLysLysPhePhe	300
Db	636144	TATGAGCGCTTTATGATTACATCTGCAATTCATTCAAAAAATTAAGAAATTAAGAAAAATTA	636085
QY	301	SerLysAlaLeuLysGlu	313
Db	636084	GAACAAATATTTAAAGAAATATATAAAGTAAACCCGACTTGCTTAAAAAACTTCCCT	636025
QY	314	AspLeuPheGluAsnLysIleGlyGlu	323
Db	636024	GAATATTTATTAAT	635968
QY	324	ThrValAspPhePheAsnLysGluIleIleArgAspProSer	342
Db	635967	ACAGAGCTTTTACCAAGTTGAAGAGATTGATTCCAATATTAACAAAAATATTAAGAAAA	635908
QY	343	ValSerAsnAspAlaLysAspLeuPheGluAsn	355
Db	635907	TTG	635851
QY	356	GlyGlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIle	375
Db	635850	TCAAAAGAAATTTTGAATAATAGCAAAATGAAT	635812
QY	376	ArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGlnGlyThrVal	395
Db	635811	TTTAAACACTATTTTAAATTAATTTCAATATAGAGATTCTAAATTAATGAATGAATACACA	635752
QY	396	AspPheIle	408
Db	635751	CAATTTATTTAGCAATTAACCTTCACTTAGCTAACAATCAGTGGGAGAA	635698
QY	409	LeuIleArgLysValIrrThrGluAlaAsp	425
Db	635697	ATAGTGCCTTTTGAAGACGAAGATATCTTTGGTGAATTAATTC	635650
QY	426	GlyGlnGlyThrValAspPheIleAsnLysGluIleArgAspProSerLysAlaLeuIle	445
Db	635649	TTAAAGATTTTCATTTCCAAAATAAAGCAAAATTAATTAAGAGACTCTGT	635599
QY	446	ArgLysValSerThrGluAlaAspAsnAsnLeuGlu	457
Db	635598	ATTGCTCCATCAAAATCTCTTTAAAGCTAAACAAATGATTGA	635554
RESULT 14			
ID	AAAI5186	standard; DNA: 1165 BP.	
AC	AAAI5186;		
XX	04-SEP-2000	(first entry)	
DT			
XX	DNA encoding	Escherichia coli	virulence proteins.
DE			
XX	Virulence protein; tafa; tab; tatc; tate; mdog; crec; regc; yggn;		
KW	ecoli; ltrdb; ltrcc; ltrce; mld2; msl; vaccine; infection;		
KW	Grim negative bacterium; ss.		
XX	Escherichia coli.		
OS			
XX			
FH	key	location/Qualifiers	
FT	CDS	2..1099	
FT		/*tag= a	
FT		/product= "virulence protein"	

FT		/note= "encodes AAY93237; no termination codon given"
FT	CDS	1102..1488
FT		/*tag= b
FT		/product= "virulence protein"
FT		/note= "encodes AAY93238"
FT	CDS	1573..1896
FT		/*tag= c
FT		/product= "virulence protein"
FT		/note= "encodes AAY93239"
FT	CDS	1939..2196
FT		/*tag= d
FT		/product= "virulence protein"
FT		/note= "encodes AAY93240; no termination codon given"
FT	CDS	2198..2533
FT		/*tag= e
FT		/product= "virulence protein"
FT		/note= "encodes AAY93241"
FT	CDS	2613..3041
FT		/*tag= f
FT		/product= "virulence protein"
FT		/note= "encodes AAY93242; no termination codon given"
FT	CDS	3051..3410
FT		/*tag= g
FT		/product= "virulence protein"
FT		/note= "encodes AAY93243"
FT	CDS	3460..3705
FT		/*tag= h
FT		/product= "virulence protein"
FT		/note= "encodes AAY93244"
FT	CDS	3791..4837
FT		/*tag= i
FT		/product= "virulence protein"
FT		/note= "encodes AAY93245"
FT	CDS	4878..7802
FT		/*tag= j
FT		/product= "virulence protein"
FT		/note= "encodes AAY93246"
FT	CDS	7816..9483
FT		/*tag= k
FT		/product= "virulence protein"
FT		/note= "encodes AAY93247"
FT	CDS	9836..10084
FT		/*tag= l
FT		/product= "virulence protein"
FT		/note= "encodes AAY93248"
FT	CDS	10134..10430
FT		/*tag= m
FT		/product= "virulence protein"
FT		/note= "encodes AAY93249"
FT	CDS	10459..10779
FT		/*tag= n
FT		/product= "virulence protein"
FT		/note= "encodes AAY93250"
XX		
PN		WO200028038-A2.
XX		
PD		18-MAY-2000.
XX		
PF		99MO-GB03721.
XX		
PR		98GB-0024569.
PR	09-NOV-1998;	98GB-0024570.
PR	17-DEC-1998;	98GB-0027814.
PR	17-DEC-1998;	98GB-0027815.
PR	17-DEC-1998;	98GB-0027816.
PR	17-DEC-1998;	98GB-0027818.
PR	13-JAN-1999;	99GB-0000708.
PR	13-JAN-1999;	99GB-0000710.
PR	13-JAN-1999;	99GB-0000711.
PR	28-JAN-1999;	99GB-0001915.
XX		
PA	(MICR-) MICROSCIENCE LTD.	
XX		

PI Crooke HR, Clarke EE, Everest PH, Dougan G, Holden DW, Shea JE;
PI Feldman RG;
XX
DR WPI: 2000-376550/32.
DR P-PSDB: AAY93237, AAY93238, AAY93239, AAY93240, AAY93241, AAY93242,
DR AAY93243.
XX
PT Peptide encoded by an operon including genes from *Escherichia coli* for
PT screening potential drugs, detecting virulence and treating conditions
PT associated with infection by a Gram negative bacterium -
XX
XX
PS Disclosure: Page 83-101; 122pp; English.
XX
CC The present sequence encodes *Escherichia coli* virulence proteins
CC The specification describes virulence proteins which are encoded
CC by an operon including *tatA*, *tatB*, *tatC*, *tatE*, *mdgG*, *crec*, *recG*, *yggN*,
CC *ech1*, *iroD*, *iroC*, *iroE*, *msl-16* genes obtained from *Escherichia*
CC *coli* K1. The virulence proteins and polynucleotides, and their vaccines
CC are useful for screening potential drugs, for the detection of virulence,
CC and for treating or preventing conditions associated with infection by
CC a Gram negative bacterium particularly *Escherichia coli*.
XX
SQ Sequence 11165 BP; 3592 A; 2087 C; 2458 G; 3028 T; 0 other;
XX
Alignment Scores:
Pred. No.: 0.375 Length: 11165
Score: 121.00 Matches: 89
Percent Similarity: 35.70% Conservative: 77
Best Local Similarity: 19.14% Mismatches: 143
Query Match: 5.13% Indels: 156
DB: Gaps: 22
US-09-807-459-2 (1-458) x AA15186 (1-11165)
QY 99 GlnLeuGluAlaIaPheIleuPheLysGlu-----SerAspAlaAsnProIa 115
DB 5604 CGCTTAGAAAGCTTTATTAGTTTACAAACACATGTACAGATTCACACCTCCT 5663
QY 116 AsnSerThrGluLysArgPheTrpMetArg-----PheArgArgGlyLysAsn 131
DB 5664 ATTAATAGCTGTTTACGACTATCTGGAAGATGCTGATTCAGACAAATTAACGAA 5723
QY 132 HisSerTyPheHisAspLeuValPheAsnLeuGluLysAsnValThrArgAspAla 151
DB 5724 TCATATTTTATTACACCCAGCAATTCATCTTCGACATGCACAC----- 5768
QY 152 AspAlaThrAspIleGluAsnPheAlaSerArgTyLeuTyMetAlaThrLeuTyTrp 171
DB 5769 -----ATTGAAGATG----- 5780
QY 172 LysThrTyThrAsnValAspGluPheGlyAlaSerPhePheAsnLysLeuSerPheThr 191
DB 5781 -----TTCAAATGAATTTGCT-----CAATTCAGTGGACT 5810
QY 192 ThrGlyLeuPheGly-----TrpGlyIleLysArgAla 202
DB 5811 GTTCTCTGTTATGCTGAAGACCAATTCATGTCGACTGGAAAAATATATATCA 5870
QY 203 LeuLysGln-----IleLeuArg 208
DB 5871 TTAACCCCAATCGCTCATTAATAAATTAAGCTATATTCATGCTGCTGCTGAT 5930
QY 209 SerAsnLeuProLeuAspIleGlyThrGluHisSerValSerArgLeuGlnHisIleThr 228
DB 5931 TCACGCTTACCCATGAGATGCTG-----GCCCTCTGAAGAGAAAAACA 5975
QY 229 SerSer-----TyrLysAsp-----TyrMetAsp 236
DB 5976 GATAGATATGCTGTTATATATATATGAGATTAAGAAATTTGCGCATGAGAT 6035
QY 237 ThrGlnIleProAlaLeuProLysPheAlaLysArgPheSerLeuMetVal----- 253
DB 6036 TCAGATACGATTTTCTA---AAAATAGAAAAAGACGATTTATCCGCTTCTGAATAT 6092

QY 254 -----ValGlnArgLeuLeuAlaThrVal----- 261
DB 6093 TTTTCTCATATCGACGCTTTGTTGGACCAATAGATTAAACAAAAGAAATGCTTCA 6152
QY 262 -----AlaGlyTyValAspThrProTyPtyLysIleTyTrp 274
DB 6153 TTAGTTGAAAAGCTGGGCGACAGACATTCATTGAAAATTAACCATATTAACAGTTTAA 6212
QY 275 MetLysLeuLysAsnPheMetVal----- 282
DB 6213 GAATGCTTGAATAATTTCTTCATCGAATCGCAAGATTTCTTAAGCAGATGCGAT 6272
QY 283 ---AsnArgValPheIleProThrLysLysPhePheAsnLysGlu-----IleArg 298
DB 6273 ATGCTGATTTATTTGTTGAGCAAAAGCACATAGAAATGAAGAACATGTTTATCT 6332
QY 299 GluProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeuPheGluAsn 318
DB 6333 AAAAGATCTAAACAACTAAACCTAAAGATAGATTAAAGAAAGATCTGAT----- 6386
QY 319 LysIleGlyGlnGlyThrValAspPheAsnLysGluIleArgAspProSerLysAla 338
DB 6387 -----GATTTTGTGATTAAGTATGATTAATGATTAATGATTAAT 6425
QY 339 LeuLysGluLysValSerAsnAspAlaLysAspLeuPheGluAsn-----LysIleGlyGln 357
DB 6426 GAATATTAATTAACCTATATCATATAAAACGAGAAATTTCTCCAGTACAGAAATTAACAGAC 6485
QY 358 GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 377
DB 6486 ACCATATATGATTTATGTTATATCAATAAATAAGAAACAAATAGCTATCTATAAAT 6545
QY 378 ValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe 397
DB 6546 CTACCTAATTTCTGCTGAT-----ATAAGAAACCTCTGAGTTGATTAACAAAGAG 6599
QY 398 IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValTyThrGluAla 417
DB 6600 TTATCTAAT---TTATGGATAGATATCAATAGAAAGACAAATAATCTGTTACACTA 6656
QY 418 AspAspLeuPheGluAsn-----LysIleGlyGlnGlyThrValAsp 431
DB 6657 AATGAGCTTAAGATTAACCTGATAGAAAGCTTATAGAACTGATTAATAAATATATGAT 6716
QY 432 PheIleAsn-----LysGluIleArgAsp----- 439
DB 6717 TTTCTCACTTACGGAAGACCTTGAAAGATCTTGAATCTACAAACAAAGTTACTATGAA 6776
QY 440 -----ProSerLysAlaLeuIleArgLysValSerThrGlu 451
DB 6777 AAAGACTTAACAAATGTTATATATGACGCTTAATAATGCTTGAAGATGCAATCTTAA 6836
QY 452 AlaAspAsnLeuLeu 456
DB 6837 GCAAAATGAGTTAAT 6851
RESULT 15
AAV71915
ID AAV71915 standard; DNA: 3883 BP.
AC AAV71915;
XX 11-FEB-1999 (first entry)
DE S. cerevisiae CIN8 DNA sequence.
XX
XX TIR1: recombinant; research; epitope mapping; modulating; CKI; yeast;
XX casein kinase I; cell growth; CIN8; cancer; viral infection; ss.
OS Saccharomyces cerevisiae.
XX
XX US5846764-A.

QY 381 GLYALGluASP-----LeuphegluAsnLysIleGlyGlnGlyThrValAspPheIle 398
DB 2551 ATGCACAGGAGAAATTTCTTCAAGAGACTAATATC---CAACCAATCTTGATGATGAC 2607
QY 399 AAsnAsnGluIleArgAspProSerIleAlaLeu----- 409
DB 2608 AAAAATGAACTGACTCTTATGAGAACCATGCAAGAAAAGCTGACTAATGTACAA 2667
QY 410 -----IleArgLysValIleThrGluAlaAspAspLeuPheGlu-----AsnLys 424
DB 2668 GACTGTGTGAAGAAATTTTAAACGAATCTCTAAATCTTCAAGCTGTATTGAGAA 2727
QY 425 IleGlyGlnGlyThrValAspPhe-----IleAsnLysGluIleArgAspProSerLys 442
DB 2728 ATCGACATATATAGACTAGATTTCACAAAATTTTATATAAAATATAGCCGAGAT----- 2781
QY 443 AlaLeuIleArgLysValSerThrGluAlaAspAsnLeu 455
DB 2782 -----CTTTCGATATATGACGAGAAATTAACACATG 2814

RESULT 16

AAH78010 standard; DNA; 3884 BP.
AAH78010;
13-NOV-2001 (first entry)
Nucleotide sequence of bImc homologue, cIn8.
bImc: kinesin related protein; fungal viability; antifungal; cIn8;
fungal infection; ss.
Saccharomyces cerevisiae.

Key Location/Qualifiers
CDS 491..3607
FT /*tag= a
FT /product= "cIn8"

US6284480-B1.

04-SEP-2001.

03-APR-2000; 2000US-0541782.

03-APR-2000; 2000US-0541782.

(CYTO-) CYTOKINETICS INC.

Nislow CE, Sakowicz R, Beraud C;

WPI: 2001-540724/60.

P-PSDB: AAG67416.

Identifying a modulator, e.g. antifungal agent, of a target protein comprising bImc or its fragment by determining enzymatic activity of a reaction, in the presence and absence of the compound, that uses ADP or phosphate produced by bImc -

Disclosure: Fig 3A-B; 47pp; English.

The present sequence encodes a bImc homologue, designated cIn8. bImc is a kinesin related protein, which is essential for fungal viability. The specification describes a method of identifying modulators of bImc. The method comprises adding a test agent to a mixture comprising bImc protein that directly or indirectly produces ADP or phosphate, subjecting the mixture to an enzymatic reaction that uses the ADP or phosphate, and determining the enzymatic activity in presence and absence of test compound. A change in the activity level between the presence and absence of the candidate agent indicates a modulator of the target protein function. The method is useful for identifying a modulator, e.g. antifungal agents, of bImc. The modulators can be used, for example, to

CC inhibit the growth or spread of fungi, mould, fruit flies, etc.. The
CC modulators can be used for preventing and treating infections caused
CC by chytridiomycetes, Hyphochytridiomycetes, Plasmidiomycetes,
CC Oomycetes, Zygomycetes, Ascomycetes, and Basidiomycetes.

Sequence 3884 BP; 1496 A; 652 C; 680 G; 1056 T; 0 other;

Alignment Scores:

pred. No.:	0.115	Length:	3884
Score:	120.00	Matches:	121
Percent Similarity:	34.75%	Conservative:	92
Best Local Similarity:	19.74%	Mismatches:	214
Query Match:	5.09%	Indels:	186
DB:	22	Gaps:	29

US-09-807-459-2 (1-458) x AAH78010 (1-3884)

QY 4 SerAspSerValGlyAspValThrLysThrLeuLeuAlaIleSerGluSerValAspSer 23
DB 1052 AGCGATGACAGCAGCAATTTATACGAGGGTTCTTTGAGTGTTCACACATTGGAACTA 1111
QY 24 AlaAlaAsnAlaTyrMetIle-----AsnSerAspMetSer 35
DB 1112 CAACAGACGATTCGTAATAAATGTCTCATTTGAACCTACACAGCAATTTGAAG 1171
QY 36 AspTyrLeuSerAlaValSerAsp-----AspPheAla 46
DB 1172 GACCTCTTGACAGCAATATGACCAAGCGCTCTAGTAATAGCGCTTGACGCCCAATTTANG 1231
QY 47 GluArgIleCysSerGlnValProLysGlySerAsnCysSerAlaSerValSerAlaTyr 66
DB 1232 AAAAATTTGGAGATTTTTCCTTCACACAGCAAAATATACCACTGACATGCTAGT 1291
QY 67 MetSerArgCysAlaLysGlnAspCys-----LeuThrLeuGln 79
DB 1292 AGTTCCAGAGTAATTCGTAGGAACTCTCCAGGCTCATTAATGATCTAACACCTTAA 1351
QY 80 Ser-----LeuLysTyrProLeuGluAlaLysTyrGlnProLeuThrLeuProAspPro 97
DB 1352 GCTGCTCTATTAGAAAGGTTTAAAGGACAAATTCACGCCGATATCCATCAAGCAACAG 1411
QY 98 TyrGlnLeuGluAlaAlaPheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSer 117
DB 1412 TATCAACACACACAGCAGATGATTCAGGACACACACTCTCTGCTGCTGCTAC 1471
QY 118 ThrGlu-----LysArgPheThrPheMetAlaPhe 126
DB 1472 ACTAATATGCTTCTAGTAACACCAACACAAATTAACGATTCGATGGCTCA 1531
QY 127 ArgArgGlyLysAsnHisSerTyrPheHisAspLeuVal----- 139
DB 1532 AATGACCAACTAATGCTATATACATCCAGATTTGCAAGATTTTCACATACAAATTCCT 1591
QY 140 -----PheAsnLeuLeuGluLysAsnValThr---ArgAspAlaAspAlaThrAsp 155
DB 1592 ATGAGAGGGGTAAACCTATATACAAAAGGCTTAAGCATGACAGAGGCTCCACTTAA 1651
QY 156 IleGluAsnPheAlaSerArg-----TyrLeuTyrMetAlaThrLeuTyrTyrThr 173
DB 1652 ATGAACGATTTTCCAGTAGATCTCATACATTTTACAATCTTGTATAGAAGCAT 1711
QY 174 Tyr-----ThrAsn 176
DB 1712 CAGGATGAACATATTAGAAATTTCCAAATGAATCTTGATTTAGCTGTTACAGAAAC 1771
QY 177 ValAspGluPheGly-----AlaSerPheAsnLysLeu 188
DB 1772 ATCAACAGATCCGAGCATTAATTAACAGTGCACAAAGAGCTGTTCAATCAACCAAACT 1831
QY 189 SerPheThrThrGly----- 193
DB 1832 CTATTGACGCTGGCAGGCTCATAAAGCACCTCGTAGATAAAGCGGCATATACCTTTC 1891

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Oy 194 -----LeuphegltYrpglYlelYsArghA 202
Db 1892 CGTGAATCGAAATGACCGCGCTTCAAGATTCCTGGTGGTAATACGAAACCGCA 1951
Oy 203 LeuYsGltInlelleAgSerAsnLeuProleuAspIleGlyThrGluHisSerValSer 222
Db 1952 CTA-----ATTGGTACTATATCCCTGCAAGAGTAACCTTGAGAAACCTGGCGT 2002
Oy 223 ArgLeuGlnHisIleThrSerSerThrLysAspTyrMetAspThrGlnIleProAlaLeu 242
Db 2003 ACATTAGAGTAT---GCTTCAAGGCTAAACATTAAGAACAG-----CCGCACTG 2053
Oy 243 ProlYsPheAlaLysArgPheSerLeuMet-----Val 253
Db 2054 GGTTCATTATTAATGAAGATATTTGGTTAAATATATACATAGAAATAGCAAAAGAT 2113
Oy 254 ValGlnArgLeuAlaThrValAlaGly-----TyrValAspThrProTyr 270
Db 2114 AATATCCATTTACTCTCTACAAAGTCCAAAGAGAAATATATAGCCAAAGATCACACTAC 2173
Oy 271 LysLysTrpTyrMetLysLeuLysAsnPheMetValAsnArgValPheIleProThrLys 290
Db 2174 AAAAATTTGAACACTGATTTTGAAGATTATAA---AATGAGTT----- 2215
Oy 291 LysPhePheAsnLysGluIleArgLysProSerLysAlaLeuLysGluLysValSerThr 310
Db 2216 -----CAAGATGTAAAGAGAAATGAAAGTTTGAATCGAATCGAAATGATTC 2263
Oy 311 AspThrLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheAsnLys 330
Db 2264 CTAGTAAAGATTAATGAACTCAAA-----GAACTATTCATTCACAAATTTGC 2314
Oy 331 GluIleArgAspProSerLysAlaLeu-----LysGlu 341
Db 2315 CAATAGATCATTTGAAACACACATGATCATTTAAGGACACAACTGATTAACAGCAT 2374
Oy 342 LysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThr----- 359
Db 2375 AAAAAGTGAATTTGAATATCCGAT---TTTATTAACAAACACACAGAGTTGACTGAGGTA 2431
Oy 360 -----ValAspPhe----- 362
Db 2432 ATGCAAAATGCCCTACATGATTACAAAAAAGACAACTTGACCTTAATCAAAAGTTTCAA 2491
Oy 363 -----IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValSerThr 380
Db 2492 ATGCAATATTACTAAGAAATTAATAAATTGAATCTACACTGTTTTCATTAACACT 2551
Oy 381 GluValAsp-----LeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIle 398
Db 2552 ATGCACAGGAAAGTATTTCTTCAAGAGCTAATATC---CAACCAAAATCTGATATGATC 2608
Oy 399 AsnAsnGluIleArgAspProSerLysAlaLeu----- 409
Db 2609 AAAAATGAGTACGACTCTTATGACACATGCAAGAAAAAGCTGAACATATGACAAA 2668
Oy 410 -----IleArgLysValTyrThrGluAlaAspAspLeuPheGlu-----AsnLys 424
Db 2669 GACTGTGAGAAAGAAATTTAAAGCAATCTCTAAATCTTCAATGTTGTTATTGAGAAA 2728
Oy 425 IleGlyGlnGlyThrValAspPhe-----IleAsnLysGluIleArgAspProSerLys 442
Db 2729 ATTCACATTAATTAAGATGATTTTCCAAAAAATTTATATAAATATAGCGGAGAAAT----- 2782
Oy 443 AlaLeuIleArgLysValSerThrGluAlaAspAsnLeu 455
Db 2783 -----CTTCTGATATTAGCGAAGAAATTAACAACATG 2815

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XX 22-SEP-1994 (first entry)
DT
XX Malarial PfEMP3 epitopic fragment clone p2b1.p12-1.
DE
XX Plasmodium falciparum erythrocyte membrane protein; pfEMP3;
KW malaria; antigen; epitope; vaccine; anti-idiotype antibody; ds.
XX
OS Plasmodium falciparum (Malayan Camp strain).
XX
FH Key Location/Qualifiers
FT CDS 3..4766
FT /tag= a
FT /note= "partial coding region; does not include
FT initiation or termination codons"
XX
PN W09403604-A.
XX
XX 17-FEB-1994.
PD
XX
XX 05-AUG-1993; 93WO-US07261.
PF
XX 07-AUG-1992; 92US-0927531.
PR
XX (SCHE ) SCHERING CORP.
PA Handumetli SM, Howard RJ, Pasloske BL, Van Schravendijk MR.
PI WPI: 1994-065693/08.
XX P-PSDB: AAR46605.
DR
XX New malaria antigen, pfEMP3 - used to isolate and produce prods.
PT for use in diagnosis, therapy and prevention of malarial
PT infection
XX
PS Claim 1; Page 72-79; 79pp; English.
XX
XX The pfEMP3 malarial antigen is recognised by monoclonal antibody Mab
CC 12C11. Nucleic acid sequences encoding part of the 315kd antigen,
CC have been isolated and sequenced. pfEMP3 is encoded on chromosome 2
CC of the P. falciparum genome and is thought to be associated with knob
CC formation and structure; malarial strains carrying deletions of the
CC gene coding for pfEMP3 exhibit a knobless phenotype.
XX
SQ Sequence 4766 BP; 2404 A; 508 C; 863 G; 991 T; 0 other;
Alignment Scores:
Pred. No.: 0.212 Length: 4766
Score: 118.50 Matches: 57
Percent Similarity: 47.57% Conservative: 31
Best Local Similarity: 30.81% Mismatches: 66
Query Match: 5.02% Indels: 31
DB: Gaps: 14
US-09-807-459-2 (1-458) x AA070102 (1-4766)
Oy 294 AsnLysGluIleArg---GluProSerLysAlaLeuLysGluLysValSerThrAspThr 312
Db 3135 AATTAAGATTAACGAAATTAAGATCTGAAGATTTAAAGAAATGCAACATAAAT 3194
Oy 313 LysAspLeuPheGluAsnLysIleGlyGlnGly-----ThrValAspPheAsn 329
Db 3195 AAAAGATTAA---CAAAATTAAGATCTGCAAGATTTAAAGAAATGCAAGATTAAAAAT 3251
Oy 330 LysGluIleArgAspPro---SerLysAlaLeuLysGluLysValSerAsnAspAlaLys 348
Db 3252 AAAGATTAACAAATTAAGAGATCTGAAGATTTAAAGAAATGCAAGCAAAAAATTA 3311
Oy 349 AspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnAsnGluIleArg 368
Db 3312 GAATTA---CAAAATTAAGATCTGAAGATTTAAAGAA-----AATGCAAGATTAA 3362
Oy 369 AspProSerLysAlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGlu--- 387

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RESULT 17
 AA070102
 ID AA070102 standard; cdna to mRNA; 4766 BP.
 XX
 AC AA070102;

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Db 3363 -----AAATAAGATTTGCAATTAAGATCTGATGATTAAGAAATGCAAGCTA 3416
Oy 388 -----AsnLysIleGlyGlyGly-----ThrValAspPheIle 398
Db 3417 AAAAATAAGATTAAGATTAAGATCTGATGATTAAGAAATGCAAGATTAATA 3476
Oy 399 AsnAsnGluIleArgAspPro---SerLysAlaLeuIleArgLysValIleThrGluAla 417
Db 3477 AATTAAGATTAAGATTAAGATCTGATGATTAAGAAATGCAAGATTAATA 3536
Oy 418 AspAspLeuPheGluAsnLysIleGlyGlyGlyThrValAspPheIleAsnLysGluIle 437
Db 3537 GAT-----TTAAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATG 3587
Oy 438 ArgAspProSerLysAlaLeuIleArgLysValSerThrGluAlaAsp----- 453
Db 3588 AAA-----AATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3641
Oy 454 AsnLeuLeuGluLys 458
Db 3642 GAACATTAATTAATAA 3656

RESULT 18
ID AAA70095
XX AAA70095 standard; DNA; 7326 BP.
AC AAA70095;
XX
DT 07-NOV-2000 (first entry)
XX
DE Plasmodium falciparum chromosome 2 related DNA sequence SEQ ID NO:228.
XX
KW Plasmodium falciparum; chromosome 2; human malaria parasite; vaccine;
KM antimalarial; malaria; protozoacide; infection; insecticide; ds.
XX
OS Plasmodium falciparum.
XX
PN WO200025728-A2.
XX
PD 11-MAY-2000.
XX
PF 05-NOV-1999; 99WO-US26796.
XX
PR 05-NOV-1998; 98US-0107131.
XX
PA (HOFF/) HOFFMAN S.
PA (CARU/) CARUCCI D.
PA (GARD/) GARDNER M.
PA (VENT/) VENTER J C.
XX
PI Hoffman S, Carucci D, Gardner M, Venter JC;
DR WPI: 2000-365347/31.
XX
PT Proteins encoded by chromosome 2 of the human malarial parasite,
PT Plasmodium falciparum, useful as antimalarial vaccines and in the
PT diagnosis of P.falciparum infection -
XX
PS Disclosure: Page 454-456; 577pp; English.
XX
CC The present invention describes proteins and their fragments (I) encoded
CC by chromosome 2 of the human malarial parasite, Plasmodium falciparum.
CC Also described are: (I) nucleotide sequences (II) encoding (I); and (2)
CC vaccines against P. falciparum infection comprising (I) or (II).
CC (I) and (II) are useful for the development of vaccines against
CC P. falciparum infection. (I) and polyclonal antisera or a monoclonal
CC antibody raised to immunogens comprising the sequences of (I), are
CC useful in the detection of infection with P. falciparum. Furthermore,
CC (I) (especially when they are rifins or secreted or membrane proteins)
CC can aid the identification of drugs to treat or prevent P. falciparum
CC infection, or they can be used to identify drug resistance in
CC P. falciparum. Sequencing of the Plasmodium chromosome 2 and the

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CC subsequent identification of proteins encoded by it will help to expand
CC our understanding of parasite biology, a process hampered by the
CC complexity of the parasitic lifecycle, and provide new targets for
CC vaccine and drug development. Parasite resistance to drugs and mosquito
CC resistance to insecticides have led to a resurgence of malaria in many
CC parts of the world, and there is a pressing need for vaccines and new
CC drugs. AAA70078 to AAA70287 and AAB18144 to AAB18352 represent nucleotide
CC and protein sequences given in the present invention, but which are not
CC specifically mentioned within the specification.
XX
SQ Sequence 7326 BP; 3588 A; 868 C; 1321 G; 1549 T; 0 other;
XX
Alignment Scores:
Pred. No.: 0.377 Length: 7326
Score: 118.50 Matches: 58
Percent Similarity: 48.658 Conservative: 32
Best Local Similarity: 31.358 Mismatches: 64
Query Match: 5.028 Indels: 31
DB: 21 Gaps: 14

US-09-807-459-2 (1-458) x AAA70095 (1-7326)
Oy 294 AsnLysGluIleArg---GluProSerLysAlaLeuLysGluLysValSerThrAspThr 312
Db 3361 AATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3420
Oy 313 LysAspLeuPheGluAsnLysIleGlyGlyGly-----ThrValAspPheIleAsn 329
Db 3421 AAAAGATTA---CAAAATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3477
Oy 330 LysGluIleArgAspPro---SerLysAlaLeuLysGluLysValSerAsnAspAlaLys 348
Db 3478 AAAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3537
Oy 349 AspLeuPheGluAsnLysIleGlyGlyGlyThrValAspPheIleAsnAsnGluIleArg 368
Db 3538 GAATTA---CAAAATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3588
Oy 369 AspProSerLysAlaLeuIleArgLysValSerThrGluAlaGluAspPheGlu--- 387
Db 3589 -----AATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3642
Oy 388 -----AsnLysIleGlyGlyGly-----ThrValAspPheIle 398
Db 3643 AAAAATAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3702
Oy 399 AsnAsnGluIleArgAspPro---SerLysAlaLeuIleArgLysValIleThrGluAla 417
Db 3703 AATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3759
Oy 418 AspAspLeuPheGluAsnLysIleGlyGlyGlyThrValAspPheIleAsnLysGluIle 437
Db 3760 AATGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3813
Oy 438 ArgAspProSerLysAlaLeuIleArgLysValSerThrGluAlaAsp----- 453
Db 3814 AAA-----AATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTAAGATTA 3867
Oy 454 AsnLeuLeuGluLys 458
Db 3868 GAACATTAATTAATAA 3882

RESULT 19
AAAB1514/C
ID AAAB1514 standard; DNA; 33303 BP.
XX
AC AAAB1514;
XX
DT 04-DEC-2000 (first entry)
XX
DE N. meningitidis partial DNA sequence gnm_61 SEQ ID NO:61.
XX
KW Neisseria meningitidis; Neisseria gonorrhoeae; genome; immunogenic;

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KM antigen: vaccine; diagnosis; infection; antibacterial; identification;
KM Meningococcus B; MenB; ds.
OS Neisseria meningitidis.
PN WO200022430-A2.
XX
XX 20-APR-2000.
XX
XX 08-OCT-1999; 99WO-US23573.
XX
XX 09-OCT-1998; 98US-0103794.
PR 30-APR-1999; 99US-0132068.
XX
PA (CHIR) CHIRON CORP.
PI Frazer CM, Hickey E, Peterson J, Tettelein H, Venter JC;
PI Meisigman V, Galeotti C, Mora M, Ratcli G, Scarselli M, Scarlato V;
PI Rappunli R, Pizze M;
PI
DR WPI; 2000-318079/27.
XX
XX Isolated nucleotide sequences of Neisseria meningitidis which can be
PT used in the diagnosis and treatment of N. meningitidis infection and
PT other Neisserial infections, for example, N.gonorrhoea -
PS
XX Claim 7; Page 1375-1385; 1760pp; English.
XX
XX The present invention describes methods of obtaining immunogenic
CC proteins from Neisseria genomic sequences. AAA81453 to AAA82414
CC represent specifically claimed Neisseria meningitidis genomic DNA
CC sequences: AAA81260 to AAA81303 and AAB25620 to AAB25663 represent
CC Neisseria DNA sequences and their corresponding proteins: AAA81254 to
CC AAA81259 and AAA81304 to AAA81321 represent PCR primers used in the
CC isolation of Neisseria meningitidis DNA sequences; and AAA81322 to
CC AAA81352 represent Neisseria meningitidis MenB polynucleotide ORF
CC sequences, which are all used in the exemplification of the present
CC invention. The nucleic acid sequences, protein sequences, and antibodies
CC against them, can be used in the manufacture of a composition. The
CC composition can be used as a medicament (or in the manufacture of a
CC medicament) for treating, preventing or diagnosing infection due to
CC Neisserial bacteria. For example, some of the identified proteins could
CC be components of vaccines against Meningococcus B; against all serotypes;
CC and/or against all pathogenic Neisseriae. Identification of sequences
CC from the bacterium will also facilitate production of biological probes,
CC particularly organism-specific probes. Attempts to make efficacious
CC Meningococcus B vaccines have failed mainly due to antigen tolerance.
CC Multivalent vaccines have also been tried but none have successfully
CC overcome antigenic variability. The provision of further, complete
CC sequences may provide an opportunity to identify secreted or surface
CC exposed proteins that may be presumed targets for the immune system and
CC which are not antigenically variable or at least more conserved than
CC other more variable regions.
XX
SQ Sequence 33303 BP; 7919 A; 8691 C; 8422 G; 8266 T; 5 other:
SO
Alignment Scores:
Pred. No.: 4 Length: 33303
Score: 117.00 Matches: 81
Percent Similarity: 35.66% Conservative: 62
Best Local Similarity: 20.20% Mismatches: 150
Query Match: 4.96% Indels: 108
DB: 21 Gaps: 19
US-09-807-459-2 (1-458) x AAA81514 (1-33303)
OY 94 LeuProAspProTyrGlnLeuGluAlaAlaPheIleuPheLysGluSerAspAlaAsn 113
DB 33123 TTGGGTGACCTTATGCGCATGCTCTCCAAATTCGCTAAGAGCAAACTTAAT 33064
OY 114 ProLaaSenSerThrGluLysArgPheThrMetArgPheArgGlyLysAsnHisSer 133
DB 33063 GGTTCGAAT-----TTATGATG-----AAAAAGGT----- 33037

OY 134 TyrPheHisAspLeuValPheAsnLeuGluLysAsnValThrArgAspAlaAspAla 153
DB 33036 -----GTGCAAAACCTATAGCATGATACGTCGGTAA-----AAG 33001
OY 154 ThrAspIleGluAsnPheAlaSerArgTyrLeuTyrMetAlaThrLeuTyrTyrLysThr 173
DB 33000 ACCCGTTAGAGAAATTTGATCG-----GTTGCACGTGCAACATTCGACCA 32953
OY 174 TyrThr -----AsnValAspGluPheGlyAla 182
DB 32952 TATGGCGCTTAATTAATCAAAATAGTAGATTAACCAATTAAGTAATTCAGACA 32893
OY 183 SerPheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrGlyLysArgAla 202
DB 32892 AGTACTATAAGCCCGTTACCGATAAT-----GGCGTTCTCCAGTCAGCAGCT 32845
OY 203 LeuLysGlnIleIleArgSerAsnLeuPro-----LeuAsp 214
DB 32844 ATGTATTAGTTAATTAATCGTTCACTTCGATATGCGGATGTTATTTGGCATTAAGCT 32785
OY 215 IleGlyThrGlu-----HisSerValSerArgLeuGlnHisIleThrSerSer 230
DB 32784 TTGGGATAGAACCCGACATGATCCACATAGCAAGCAGTAATATCCAGACGATAC 32725
OY 231 TyrLysAspTyrMetAspThrGlnIleProAlaLeuProLys----- 244
DB 32724 GAAGGCGATATAGAACACATTAATATCTGCTTTAGTAATTAAGATTTGATGATCTTTT 32665
OY 245 PheAlaLysArgPheSerLeuMetValAlaGlnArgLeuAlaThrValAlaGly--- 263
DB 32664 AAAGAGAACGCAATTTACTTTTTCATCATCTGTGATGATGATGTAACAAAGTGGTTT 32605
OY 264 ---TyrValAspThrProTyrTyrLys-----LysTrp-----TyrMetLys 276
DB 32604 GATATTACAATATAGATGCTTGGCAAAAATTTGGAGCTTGGGTAATGGATATACATAT 32545
OY 277 LeuLysAsnPheMetValAsnArgValPheIleProThrLysLysPheAsnLysGlu 296
DB 32544 TTATATAAAGCTGTGTAAAGAGAGAGCTGATGAAATATTGATGCTTAATTAATTAAC 32485
OY 297 IleArgGluProSerLysAlaLeuLysGluLysValSer----- 309
DB 32484 ATCAAGCAAGGAATGAAGCTTTTAAATGAATCAATACCTGGTTCATGATATGAAA 32425
OY 310 -----ThrAspThr 312
DB 32424 GCTGCTGGCAAGAAATTTGAGATGACTTAATATACAGCTGAATATATCTCAGCTGAGCT 32365
OY 313 LysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIle 332
DB 32364 GCCCAATTAATCTAATAAGCATAGTACTAGCAATACTCAAGAAATAGAAAAGCTGTC 32305
OY 333 ArgAspProSerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspPheGlu 352
DB 32304 AAA---GCCATTAAAGAAATTTGCTGTAATAAATAATGCTGCTCCGATTTGGCTGAC 32248
OY 353 AsnLysIleGlyGlnGlyThrValAspPheIleAsnAsnGlnIleArgAspProSerLys 372
DB 32247 GGTTC-----GCAGAGAAACCTAAA 32227
OY 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyLys 392
DB 32226 CAAGTAGTGAAGATTTGCTCAACCCCAAGAAAGATACGAAAT-----GCCAAA 32173
OY 393 GlyThrValAspPheIleAsnAsnGlnIleArgAspProSerLysAlaLeu-----Ile 410
DB 32172 TCCACAGCCGGAAGAGCTGCTCAACACCTGAGAAATTTTAAAGCGTTTGGCCAGTTT 32113
OY 411 ArgLysValTyrThrGluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrVal 430
DB 32112 AAAGATCTGGCGGAATAATTTAGAGATCTGTTCCCAATTCG-----GAAGGCTGATC 32059

Oy 431 Asp 431
 |||
 Db 32058 GAT 32056
 RESULT 20
 ID AAF21610 standard; DNA: 349980 BP.
 AC AAF21610;
 DT 13-MAR-2001 (first entry)
 DE Neisseria meningitidis B nucleotide sequence SEQ ID NO:111.
 XX
 XX Neisseria meningitidis; Neisseria gonorrhoeae; immunogenic; vaccine;
 KW diagnosis; antigen; detection; infection; gene therapy; antibacterial;
 ds.
 XX
 OS Neisseria meningitidis.
 XX
 XX WO200066791-A1.
 PN
 PD 09-NOV-2000.
 XX
 PF 08-MAR-2000; 2000WO-US05928.
 PR 30-APR-1999; 99US-0132068.
 PR 08-OCT-1999; 99WO-US23573.
 PR 28-FEB-2000; 2000GB-0004695.
 XX
 PA (CHIR) CHIRON CORP.
 PA (GENO-) INST GENOMIC RES.
 XX
 XX Pizsa M, Hickey E, Peterson J, Tettelin H, Venter JC, Maignani V;
 PI Galeotti C, Mora M, Ratti G, Scarselli M, Scariato V, Rappunli R;
 PI Frazer CM, Grandi G;
 XX
 DR WPI: 2000-647603/62.
 XX
 XX Neisseria meningitidis B full length genome sequence and open reading
 PT frames are used to detect, treat and prevent Neisserial infections -
 PS
 PS Claim 7; Appendix A: 692pp; English.
 XX
 CC The present invention describes the full length genome of
 CC Neisseria meningitidis B (NMB). The sequences in AAF21544 and AAF21607
 CC to AAF21613 represent fragments of the NMB genomic sequence, as the
 CC sequence was too long to go in a record on its own it was split into 8
 CC sequences which overlap each other at the beginning and end of each
 CC sequence by 49980 bp (i.e. the last 49980 bp of AAF21544 is repeated at
 CC the beginning of AAF21607, the last 49980 bp of AAF21607 are repeated at
 CC the beginning of AAF21608, and so on). AAF21545 to AAF21588 encode the
 CC Neisseria proteins given in AAB58550 to AAB58593, and AAF21589 to
 CC AAF21606 represent PCR primers which are used in the exemplification of
 CC the present invention. The NMB genome and fragments from it have
 CC antibacterial activity, and can be used in vaccines and gene therapy.
 CC Neisseria nucleic acids, proteins and/or antibodies which binds to the
 CC proteins can be used in compositions for treating or preventing infection
 CC due to Neisserial bacteria or as a diagnostic reagent for detecting the
 CC presence of Neisserial bacteria or of antibodies raised to Neisserial
 CC bacteria. Computers, computer memory, computer storage medium or computer
 CC databases can be used in a search to identify open reading frames (ORFs)
 CC or coding sequences within the NMB genome. The DNA sequences provide
 CC further opportunities to find antigenic or immunogenic proteins which are
 CC more effective in vaccines than the outer membrane proteins currently
 CC used.
 CC
 XX Sequence 349980 BP; 86771 A; 92803 C; 86340 G; 84066 T; 0 other;
 Alignment Scores:
 Pred. NO.: 92.3 Length: 349980
 Score: 117.00 Matches: 81
 Percent Similarity: 35.668 Conservative: 62

Best Local Similarity: 20.20%		Mismatches: 150	
Query Match: 4.96%		Indels: 108	
DB:	21	Gaps:	19
US-09-807-459-2 (1-458) x AAF21610 (1-349980)			
OY	94	LeuProAspProTyrGlnLeuGluAlaIalaPheIleuPheLysGluSerAspAlaAsn	113
Db	243690	TTGGGTGACCGTGTATGGCGATGATGCTGCTCAATTTGCGTAAGGATGGCAACTTAAT	243749
OY	114	ProIaAsnSerThrGluLysArgPheThrPheTargPheArgGlyLysAsnHisSer	133
Db	243750	GGTTTGAT-----TTATGGATG-----AAAAAGGT	243776
OY	134	TyrPheHisAspLeuValPheAsnLeuGluLysAsnValThrArgAspAlaAspAla	153
		-----GTTGAAACCCTTGGATGATACGGTCGGTAA-----AAG	243812
Db	243777	-----	-----
OY	154	ThrAspIleGluAsnPheIaSerArgTyrLeuTyrMetIaThrLeuTyrLysThr	173
Db	243813	ACCCGTTTGGAAATTTGATCG-----GTTCCACTGCACACTTTCCAGCAA	243860
OY	174	TyrThr-----AsnValAspGluPheGlyAla	182
Db	243861	TATCGCGCTCAATTAATCAAAATATATGCTAGATTTACCCAACTACTAGTGAATTTAGACAG	243920
OY	183	SerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrGlyLysArgAla	202
Db	243921	AGTTACTATTAAGCCGTTACCGATAAT-----GGCGTTTCTCCAGTCAGCT	243968
OY	203	LeuLysGlnIleIaArgSerAsnLeuPro-----LeuAsp	214
Db	243969	ATTGATTTAGTTAATTAATTCCTTCCGATATGCGGATGCGGATTTTGGCATTAAGT	244028
OY	215	IleGlyThrGlu-----HisSerValSerArgLeuGlnHisIleThrSerSer	230
Db	244029	TTGGGATAGAGAACCCGAACGATCTACCAATAGACAGACGATAATTAATCCGAACGGTAGC	244088
OY	231	TyrLysAspTyrMetAspThrGlnIleProAlaLeuProLys-----	244
Db	244089	GAAAGGCTAATATAGAAAGCACTTAATATCTGCTTTAGATTAAGCATTTGATGATCTTTT	244148
OY	245	PheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThrValAlaGly---	263
Db	244149	AAAGAGAGCATTTTACTTTTTCACATCTGTGATGATGATGAACAAAGTTAGGTGT	244208
OY	264	---TyrValAspThrProTyrLys-----LysTrp-----TyrMetLys	276
Db	244209	GAATATACATATAGATGCTTGGCAAAAATTTGAGCGTTGGGTAATGGGATATCATGAT	244268
OY	277	LeuLysAsnPheMetValAsnArgValPheIleProThrLysLysPhePheAsnLysGlu	296
Db	244269	TTATATATAAAGTGTGTATATAAAGAGAGAGCTGCAGCTGGATATTTCGATCGCTTAATTAAC	244328
OY	297	IleArgGluProSerLysAlaLeuLysGluLysValSer-----	309
Db	244329	ATCAAGCAAGAAATGAAGCTTTTAAATAATGAATCAATAGCTTGCTCATGATATGAAA	244388
OY	310	-----ThrAspThr	312
Db	244389	GCTGCTGGCAAGAAATTTGGAGATGACTTAATATACACAGTGGAATATCTCACTCAAGCT	244448
OY	313	LysAspLeuPheGluAsnLysIleGlyGlnIleThrValAspPhePheAsnLysGluIle	332
Db	244449	GCCGAATTAATCTATATGACATGTAGACATATCTAGTCAAGAAATAGAAAAAAGGTCTC	244508
OY	333	ArgAspProSerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGlu	352
Db	244509	AAA---GCCATTAATAAGATGTGCTGGAATAAATGCTGCTCCGATTTGGCTGAC	244565
OY	353	AsnLysIleGlyGlnIleThrValAspPheIleAsnGlnIleArgAspProSerLys	372
Db	244566	GCTTCA-----GCAGGAAGAGCTAA	244586

CC primers or probes to isolate homologous sequences in other plants
 CC (see also AAV84692-93).

XX Sequence 10478 BP; 3195 A; 1770 C; 1986 G; 3527 T; 0 other;

Alignment Scores:

Pred. No.:	1.89	Length:	10478
Score:	113.50	Matches:	99
Percent Similarity:	38.29%	Conservative:	89
Best Local Similarity:	20.16%	Mismatches:	190
Query Match:	4.81%	Indels:	113
DB:	20	Gaps:	27

US-09-807-459-2 (1-458) x AAV84691 (1-10478)

```

OY 14 LeuLeuAlaIaSerGluSerValAspSerAlaIaAsnAlaIaTyrMetIleAsnSerAsp 33
    :::::|||||  |||  |||  :::::  :::::  :::::  :::::  :::::  :::::  :::::
Db 8178 ATCTTGCAGAAATTCGAGAGAGAGAGAGATGATCTGACTATCATCACTGCTCAGCGAT 8237
OY 34 MetSerAspTyrLeuSerAlaValSerAspAsnPheAlaGluArgIle----- 49
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8238 CTCAATCTCTACTTGATACAGACATGATGAAATTCATCATCGATTCTACGGTAGCTC 8297
OY 50 -----Cys-----SeriGlnValPro 54
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8298 TCCTCAATGCGCTTCTGCTTCTCCGCTGTAATCTCCGCCGCCATGATTCCTCCGGTTGA 8357
OY 55 LysGlySerAsnCySserAlaSerValSerAlaTyrMetSerArgCysAlaGlnAsp 74
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8358 TCAGCCCTCTGCTATTTTATTTGATGCGCCAGCGATATCTCAGCTCTCTTCCACGAT 8417
OY 75 CysLeuThrIleGlnSerLeuIleTyrProLeuGlnAlaIaTyrGlnProLeuThrIleu 94
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8418 TGC-----CCCGTTTCACTTCTCTCTCTCCGATTGCGTTT 8453
OY 95 ProAspProTyrGlnLeuGlnAlaIaIaPheIleuPheIleuPheIleuSerAspAlaAspPro 114
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8454 CTTCCTACTTGGCTCATTCGCCGCTACCTCAAAACGCCGATGATTCGCCGCTCA 8513
OY 115 AlaAsnSerThrGlnIleuArgPheThrMetArgPheArgArgGlyLysAsnHisSerIle 134
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8514 GCGAGTAATTCGACAGAGAGAGAGAGAAAGAAAGAGAGAGAGAGAGAGAGAGAGAGAG 8564
OY 135 PheHisAspLeuValPheAsnLeuGlnIleuLysAsnVal-----ThrArgAspAlaAspAla 153
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8565 -----AAGAGCATTTAGCGTATGACATGACATGCGGAGAG 8597
OY 154 ThrAspIleGluAsnPheAlaSerArgTyrIleuTyrMetAlaThrLeuTyrTyrIleSerThr 173
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8598 ACTGAAAGAGGTGATTTGATGCGCAAAATGATGATTATAGTG----- 8639
OY 174 TyrThrAsnValAspGluPheGlyAla-----SerPhePheAsnLysLeuSerPheThr 191
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8640 -----TTAGACAAACCTTGGCTCGGCTTCTGAGTTTGTTCATGATGATGATTTCTCT 8690
OY 192 ThrGlyLeuPheGlyTyrGlyIleLysArgAlaLeuLysGlnIleIleArgSerAsnLeu 211
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8691 GATAGTTTG-----AAATCTTTGATCAAAACCTG---AGTGAGATT 8729
OY 212 ProLeuAspIleGlyThrGluHisSer-----ValSerArgLeuGlnHisIle 227
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8720 CCTTAA---TTGGCGTGTGACGACACTGTGAGTTTGAATTATGATGATGATGAAATG 8786
OY 228 ThrSerSerIle-----LysAspTyrMetAspThrGlnIleProAla 241
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8787 TGTGGGAAGATTCTGGGAGAGAGTGTGAATCTGACCATGAGATGAGCATCACTCACTGCT 8846
OY 242 LeuPro-----LysPheAlaLysArg 248
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8847 GCTGATTTTCAAGCTTTTGACACCGTTTCTTATGCGTTTACATCAACGAGAAAT 8906
OY 249 PheSerLeuMetValValGlnArgLeuLeuAlaThrValAlaGlyTyrValSerThrPro 268
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
  
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Db 8907 TTTGGCTGGCATTTGTGTCAAGAAATGATGATTTGGCT-----AAAGATAACCT 8960
OY 269 TrpTyrLysLysTyrTyrMetLysLeuLysAsnPheMetValAsnArgValhellePro 288
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 8961 GAATTGAAAAAGTTGTCTTAATTTGCTTAAGTTTCTGCTTCATCAATGAGCA-----CCT 9014
OY 289 ThrLysLys-----PhePheAsnLysGluIleArgGluProSerLysAlaLeu 304
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 9015 GAGAAAGCCGAGACCCCGCTGATTTGACAGAGAGCGGCTGCTGAGATTATAAGCAAG 9074
OY 305 LysGluLysValSerThrAspThrLysAspLeuPheGluAsnLysIleGlyGlnGlyThr 324
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 9075 GAGTTGAGAGGCCAATCAAGAGTTTGTGAT---TTGTATGATGATGCTGCAAGGAGAG 9131
OY 325 ValAspPhePheAsnLysGluIleArgAspProSerLysAlaLeuLysLysValSer 344
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 9132 TCTAATTTTAAAGTATTAGCTGTT---GATATTATACCTCTGTGTGTAAGCTCATTAAGA 9188
OY 345 AsnAspAlaLysAspLeu-----PheGluAsnLysIleGlyGlnGlyThr 359
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 9189 AACCTCTAGAGATATTATGTTCAAGAGATGCGGTTCGAAAGATTTCGCGGCTTGGTTGT 9248
OY 360 ValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValSer 379
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 9249 ATTGATGCTTTAGTTACAGCGGCTTTCAGACAGAGC---GCTTTGATTTAGA----- 9296
OY 380 ThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsn 399
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 9297 -----GCTCAGAGCTTTG---TCCAACTTGGCTCAA---GTTGTGAGTCTTGTCT 9341
OY 400 AsnGluIleArgAspProSerLysAlaLeuIleArgLysValTyrThrGluAlaAspAsp 419
    :::::  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 9342 GGTGATGAAAG-----AGTAGTCGATCCGAAACAGCCCTTGGTTAAACGTGAG 9395
OY 420 LeuPheGluAsnLysIleGlyGlnGlyThrVal---AspPheIleAsnLysGluIleArg 438
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 9396 ACTTCAGAG-----GGAAGAGGTGCACTTAACGACCTTTTGAAGAAAGATGTGTG 9446
OY 439 AspProSerLysAlaLeuIleArgLysValSer 449
    |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 9447 GAT---GAGAGAGCGGCTGTAAAGAGAGAGAGCT 9476

RESULT 24
ID AAS76165 standard; cDNA; 4282 BP.
XX AAS76165;
AC AAS76165;
DT 13-FEB-2002 (first entry)
DE DNA encoding novel human diagnostic protein #11969.
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX Homo sapiens.
OS
XX WO200175067-A2.
XX
XX 11-OCT-2001.
XX
XX 30-MAR-2001; 2001WO-US08631.
XX
XX 31-MAR-2000; 2000US-0540217.
XX 23-AUG-2000; 2000US-0649167.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Drmanac RT, Liu C, Tang YT;
XX
XX WPT: 2001-639362/73.
XX
XX P-PSDB; ABG11978.
XX
  
```

PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX
PS Claim 1; SEQ ID NO 11969; 103bp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations in
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 4282 BP; 1735 A; 915 C; 751 G; 881 T; 0 other:
SO
Alignment Scores:
Pred. No.: 0.64 Length: 4282
Score: 113.00 Matches: 109
Percent Similarity: 34.04% Conservative: 52
Best Local Similarity: 23.04% Mismatches: 166
Query Match: 4.79% Indels: 146
DB: 23 Gaps: 24
US-09-807-459-2 (1-458) x AAS76165 (1-4282)
QY 55 LysGlySerAsnCySerAlaSerValSerAlaIleuMetSerArgCysAlaIleuGlnAsp 74
DB 1725 AAGGGATTTCAATTTAGCAAAAGAGAGAAATGTCCTGTTCCAGCAGCATGAT 1784
QY 75 CysLeuThrLeuGln-----SerLeuTyrProLeuGlnAlaIleu----- 88
DB 1785 TGTTCATCTAGAAAACCCCATCGTCCTCAGCCCAAAATCTCTTAAGCTGATACCACTT 1844
QY 89 TyrGlnProLeuThrLeuProAspPro-TyrGlnLeuGlnAlaIleuPheIleuPheIleu 108
DB 1845 CAGCAAGTCTCAGGATCAAAATCATGTACAAAAATCAACAGCATCTCTTATAC----- 1899
QY 108 sGluSerAspAlaAsnProAlaAsnSerThrGlnIleuArgPheThrPheIleuArg 128
DB 1900 -----ACCAATACACAGACAAA----- 1917
QY 128 gGlyAsnHisSerTyrPheHisAspLeuValPheAsnLeuLeuGlnIleuAsnVal-- 147
DB 1918 -----CAAGAGAGCAATTCATGAGTGAACCTCCATTCAACAATCTCCAAAGAGAAATAAA 1973
QY 148 -----ThrArgAspAlaAspAlaThrAspIleuGlnIleuAsnPheIleu 161
DB 1974 ATACCTAGATCCAACTTACAAAGGATGTGAAGACCTCTTCAAGAGAACTTCAAAAC 2033
QY 161 r----- 161
DB 2034 ACTGCTCAATGAATAAAGAGATCAAAACAAATGCAAGAACCTTCATGCTTATGAGGT 2093
QY 162 -----ArgTyrLeuTyrIleuValThrLeuTyrTyrLeuTyrThrAsnVal 177
DB 2094 AGGAGAGAAATCAATCTCGTGAATAATGCGCATCTCTCCCAAGATATTTAC----- 2145

QY 177 LAspGluPheGlyVal-----SerPheAsnIleuSerPhe 190
DB 2146 -----AGATTCAATCCATCCCATCAAGCTACCAATCACTTCTTACAGAAATGGAAAA 2201
QY 190 eThrThrGlyLeuPheGlyTyrPglyIleuArgAlaLeuIleuIleuIleuArgSerAs 210
DB 2202 AACTACTTTAAAGTTCAATTCGAAACAAAGAGACCCCTC-----ATTGCCAAGTCATC 2255
QY 210 nLeu--ProLeuAspIleuGlyThrGlnHisSerValSerArgLeuGlnHisIleuThrSer 229
DB 2256 CCTTAGCGCAAGAAAGAAACAAATCGAGGCAATCAGCTACCTACCTGACTTCAAACTATCA 2315
QY 230 SerTyrIleuAspTyrMetAspThrGlnIleProAlaLeuProIleuPheAlaIleuArgPhe 249
DB 2316 GCGTACAGTAAAC-----CAAAACATCATGGTACTGCTACCAAAA-----CAAGATAT 2363
QY 250 SerLeuMetValValGlnArgLeu-----AlaThr-ValAlaGlyTyrVal-- 265
DB 2364 AGATCAATGCAAGAACAGACAGCCCTCAGAAATATATSCCATATATCAACTATTCGAT 2423
QY 266 -----AspThrPro-----Trp-----TyrIleuSerTrp-- 273
DB 2424 CTTTGACAAACCTGACAAACAAAGCAATGGGAAAGATTCCCTATTATTAATATGATG 2483
QY 274 -----TyrMetIleuIleuAsnPheMetValAsnAr 284
DB 2484 CTGGAAACACTGCTAGCATATGTAGAAAGCTAAACCTGATCCCTTACGCTTA 2543
QY 284 gValPheIleuProThrIleuIleuSerPhePheAsnIleuGlnIleuArgGluProSer--LysAl 303
DB 2544 TACAAAATTTGATTCATCAAGATGATTAAGAACTTACATGTAGACCTTAACCATTAAC 2603
QY 303 aLeuIleuGlnIleuValSerThrAspThrIleuAspLeuPheGlnAsnIleuGlnIleu 323
DB 2604 CCTAGAGAAACAACTAGGCAATACCATTCACAGAC-----ATAGGCAATGGG 2648
QY 323 yThrValAspPhePheAsnIleuIleuArgAspProSerIleuAlaLeuIleuGlnIleuVal 343
DB 2649 CAAAG-----GACTTCATGTCTTAA-----ACACAAACAAACCAATGGCAACAAAGC 2693
QY 343 LSerAsnAspAlaIleuAspLeuPheGlnAsnIleuGlnIleuThrValAspPheIleu 363
DB 2694 CAAATTTGCAAAATGGATCTAATTAATACTAAAG----- 2727
QY 363 eAsnAsnGlnIleuIleuArgAspProSerIleuAlaLeuIleuArg-----LysValSerThrGln 381
DB 2728 -----AGCTTCGCGACAGCAAGAAAGAACTACCATCAGATGAGTGAACGCAACTACAC 2780
QY 381 yAlaGluAspLeuPheGlnIleuIleuGlnIleuGlnIleuThrValAspPheIleuAsnGln 401
DB 2781 ATCGGAGAAATAATTTT-----GC 2798
QY 401 uIleArgAspProSerIleuAlaLeuIleuArgIleuValIleuThrGlnIleuAspLeuPhe 421
DB 2799 AATTTACTCATTCGACAAAGGCTAATATTCAGAGCTTACATGAACTCAAAATAATTTA 2858
QY 421 eGlnAsnIleuIleuGlnIleuGlnIleuThrValAspPheIleuAsnIleuGlnIleuArgAspPro 441
DB 2859 CAACAAAAA-----ACAAACAAACCCATCAAAAGGCGC 2894
QY 441 rLysAlaLeuIleuArgIleuValSerThrGln 451
DB 2895 AAGGATATGACAGACACTTCTCCAAAGAA 2925
RESULT 25
AAS74637 standard; cDNA; 11087 BP.
XX
XX AAS74637:
AC AAS74637:
XX
XX 13-FEB-2002 (first entry)
XX
XX DNA encoding novel human diagnostic protein #10441.

XX	AAK68992:	
AC		
DT	06-NOV-2001	(first entry)
XX		
DE	Human	Immune/haematopoietic antigen genomic sequence SEQ ID NO:23804.
XX		
KW	Human; Immune; haematopoietic; Immune/haematopoietic antigen; cancer;	
XX	cytostatic; gene therapy; vaccine; metastasis; ds.	
XX		
OS	Homo sapiens.	
PN	WO200157182-A2.	
PD		
XX	09-AUG-2001.	
PF		
XX	17-JAN-2001;	2001WO-US01354.
PR		
PR	31-JAN-2000;	2000US-0179065.
PR	04-FEB-2000;	2000US-0180628.
PR	24-FEB-2000;	2000US-0184664.
PR	02-MAR-2000;	2000US-0186350.
PR	16-MAR-2000;	2000US-0189874.
PR	17-MAR-2000;	2000US-0190076.
PR	18-APR-2000;	2000US-0198123.
PR	19-MAY-2000;	2000US-0205515.
PR	07-JUN-2000;	2000US-0209467.
PR	28-JUN-2000;	2000US-0214886.
PR	30-JUN-2000;	2000US-0215135.
PR	07-JUL-2000;	2000US-0216647.
PR	07-JUL-2000;	2000US-0216880.
PR	11-JUL-2000;	2000US-0217487.
PR	11-JUL-2000;	2000US-0217496.
PR	14-JUL-2000;	2000US-0218290.
PR	26-JUL-2000;	2000US-0220963.
PR	26-JUL-2000;	2000US-0220964.
PR	14-AUG-2000;	2000US-0224518.
PR	14-AUG-2000;	2000US-0224519.
PR	14-AUG-2000;	2000US-0225213.
PR	14-AUG-2000;	2000US-0225214.
PR	14-AUG-2000;	2000US-0225266.
PR	14-AUG-2000;	2000US-0225267.
PR	14-AUG-2000;	2000US-0225268.
PR	14-AUG-2000;	2000US-0225270.
PR	14-AUG-2000;	2000US-0225447.
PR	14-AUG-2000;	2000US-0225757.
PR	14-AUG-2000;	2000US-0225758.
PR	18-AUG-2000;	2000US-0226279.
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PR	23-AUG-2000;	2000US-0227009.
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PR	01-SEP-2000;	2000US-0229287.
PR	01-SEP-2000;	2000US-0229343.
PR	01-SEP-2000;	2000US-0229344.
PR	01-SEP-2000;	2000US-0229345.
PR	05-SEP-2000;	2000US-0229509.
PR	05-SEP-2000;	2000US-0229513.
PR	06-SEP-2000;	2000US-0230437.
PR	06-SEP-2000;	2000US-0230438.
PR	08-SEP-2000;	2000US-0231242.
PR	08-SEP-2000;	2000US-0231243.
PR	08-SEP-2000;	2000US-0231244.
PR	08-SEP-2000;	2000US-0231413.
PR	08-SEP-2000;	2000US-0231414.
PR	08-SEP-2000;	2000US-0232080.
PR	08-SEP-2000;	2000US-0232081.
PR	12-SEP-2000;	2000US-0231968.
PR	14-SEP-2000;	2000US-0232397.
PR	14-SEP-2000;	2000US-0232398.
PR	14-SEP-2000;	2000US-0232399.

PR	14-SEP-2000	2000US-02324200
PR	14-SEP-2000	2000US-0232400
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PR	20-OCT-2000	2000US-0241787
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PR	17-NOV-2000	2000US-0249245
PR	17-NOV-2000	2000US-0249246
PR	17-NOV-2000	2000US-0249264
PR	17-NOV-2000	2000US-0249265
PR	17-NOV-2000	2000US-0249297
PR	17-NOV-2000	2000US-0249299
PR	17-NOV-2000	2000US-0249300
PR	01-DEC-2000	2000US-0250160
PR	01-DEC-2000	2000US-0250191
PR	05-DEC-2000	2000US-0250130
PR	05-DEC-2000	2000US-0251988
PR	05-DEC-2000	2000US-0256719

Seq ID	Seq	Length	Score	Percent Similarity	Best Local Similarity	Query Match
06-DEC-2000	2000US-0251479	15.5	112.50	34.44%	20.39%	4.77%
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08-DEC-2000	2000US-0251989	74	74	100%	107	16
08-DEC-2000	2000US-0251990	74	74	100%	107	16
11-DEC-2000	2000US-0254097	74	74	100%	107	16
05-JAN-2001	2001US-0259678	74	74	100%	107	16
XX	(HUMA-) HUMAN GENOME SCI INC.					
XX	Rosen CA, Barash SC, Ruben SM;					
XX	WPI: 2001-483426/52.					
XX	Nucleic acids encoding human immune/hematopoietic antigen polypeptides,					
XX	useful for preventing, diagnosing and/or treating cancers and					
XX	metastasis -					
XX	Disclosure; SEQ ID NO 23804; 3071bp + Sequence Listing; English.					
XX	AAK54951 to AAK64702 encode the human immune/hematopoietic antigen (I)					
XX	amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic					
XX	activity, and can be used in gene therapy and vaccine production. (I)					
XX	treatments and polynucleotides may be used in the prevention, diagnosis and					
XX	treatment of diseases associated with inappropriate (I) expression. For					
XX	example, they may be used to treat disorders associated with decreased					
XX	expression by rectifying mutations or deletions in a patient's genome					
XX	that affect the activity of (I) by expressing inactive proteins or to					
XX	supplement the patients own production of (I). Additionally, (I)					
XX	polynucleotides may be used to produce the secreted (I), by inserting					
XX	the nucleic acids into a host cell and culturing the cell to express the					
XX	protein. (I) proteins and polynucleotides may be used to prevent, the					
XX	diagnose and treat immune/hematopoietic-related diseases, especially					
XX	cancers and cancer metastases of hematopoietic-derived cells. AAK64703					
XX	to AAK87694 represent human immune/hematopoietic antigen genomic					
XX	sequences from the present invention. AAK54942 to AAK54950 and AAM82169					
XX	represent sequences used in the exemplification of the present invention					
XX	Sequence 42738 BP; 15539 A; 8490 C; 8408 G; 10301 T; 0 other;					
XX	Alignment Scores:					
XX	Pred. NO.:	15.5	Length:	42738		
XX	Score:	112.50	Matches:	74		
XX	Percent Similarity:	34.44%	Conservative:	51		
XX	Best Local Similarity:	20.39%	Mismatches:	131		
XX	Query Match:	4.77%	Indels:	107		
XX		22	Gaps:	16		
US-09-807-459-2 (1-458) x AAK68992 (1-42738)						
0Y 174	TYRTThrsanValaSPGcIuPneqIyAlaseRPherheaSnlyseuSerPheThrly 193					
Db 40901	TACATCATATACACAAACAAAGGAGCGCAATGATGAGTAATCTCATTCACAACTGCT					
0Y 194	-----LeupneqIyTPRgIyIlelysaRgAlaleuLyscInlleIeaRg 208					
Db 40961	TCAAAGAAATAAATACCTAGAGATCCAACTTACAAAGGATGTGAAGGCGCTTCAAG 41020					
0Y 209	serIsnleu---ProIeasRPIleqIyThGInHIsseRValserIrgIeugInHIsIte 227					
Db 41021	GAGAACTTCAACACCACTGCGCACAAAGAAATAAAGAGAGATTCACAAACAAATGGAAAGACATT 41080					
0Y 228	ThIseRseRlyRlysaSPyRmetasp---ThGInIleProAlaleuProIysPheala 246					
Db 41081	CCATGCTCATGTGGTGAAGAAACAAATATTCGTAATAATGGCCATACTGCCCAAGTAATT 41140					
0Y 247	LyasRgPheSerIeueMetValValgIn-----ArgIeueula 259					
Db 41141	TATGATTCATTCGGTTCCTCCCATTAATAGCTTACCAATGACATTCTTCACAGAAATTGGAAAAA 41200					
0Y 260	ThIValaInqIyRVal----- 265					

Db	41201	ACIGCTTTAACTTCATATGAAACAAACAAAAAGAGCCTGCATTTGCCAACTCAACCTAAAGC	41260
Qy	265	-----	265
Db	41261	CAAAAGAACAAAGCTGGAGAGCATCAGCTACTGACTTCAACATATACAGAGCTACA	41320
Qy	266	-----AspThrProTyrTyrIleValStryptophyMetLys-----	276
Db	41321	CTAACCAAAACAGCATGGTAC-----TGTACCAACAAACAGATATAGACCAATGGAAC	41374
Qy	277	-----LeuLysAsnPhenMetValAsnArgValPhe	286
Db	41375	AGAACAGAGCCTCAGAAATATACACCAACATCTACCAACTATCTGATTTTGACAAACCT	41434
Qy	287	IleProthrLysLysPhe-----PheAsnLysGluIleArgGluProSer	301
Db	41435	GACAAACAAACAAATATGGGAAAGATTCCTCATTTTAATTAACGGTGGTGGAAAAACGG	41494
Qy	302	LysAlaLeuLysGluLysValSerThrAspThrLysAspLeuPheGluAsnLysIleGly	321
Db	41495	CTAGGCATATCTAGAACAGCTGAACTGGATCCCTTCCTTACACTTATACAAAAATTTAAT	41554
Qy	322	GInGlyThrValAspPheheAsnLysGluIleArgAspProSer--LysAlaLeuLys	340
Db	41555	TCAGATGGATGATTAAGACTTAAAC-----GTTAGACCTAAACCATATAAACCTTAGAA	41608
Qy	341	GluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGInGlyThrVal	360
Db	41609	GAAGAACTTGGCAATATATCTTCAGGAC-----ATAGGCATGGGAAG---	41650
Qy	361	AspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValSerThr	380
Db	41651	GACTCATCTGTCTAAA-----ACACCAAAAGCAATGGCAACAAAGGCCAAATTT	41698
Qy	381	GlyAlaGluAspLeuPheGluAsnLys-----IleGlyGInGlyThrValAspPhe	397
Db	41699	GACAAATGGATCTATTTAACTAAAGAGCTTTCGACAGCAAAAGAAATCATCCATCAGA	41758
Qy	398	IleAsnAsnGlu-----IleArgAspProSerLysAla	408
Db	41759	GTCAGACAGGCACCTACAGCAATAGGAGAAAGTTTTCGACATCTACTCAGCTGACAAAGG	41818
Qy	409	LeuIleArgLysValTyrThrGluAlaAspAspLeuPheGluAsnLysIleGlyGInGly	428
Db	41819	CTAAATATCCAGATCTACATGAACTCAACAAATTTTACAGAAAAA-----	41866
Qy	429	ThrValAspPheIleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysVal	448
Db	41867	-----ACAAACAAACCCCATCAAAAAGTGGGCAAAAGATATGACAGACACTTC	41914
Qy	449	SerThrGlu 451	
Db	41915	TCAAAAAGAA 41923	
RESULT 28			
AXX35720			
ID	AXX35720 standard; cDNA; 1422 BP.		
XX			
XX	AAAX35720;		
XX			
XX	09-JUL-1999 (first entry)		
DE			
XX	cDNA encoding a protein identified by the signal sequence trap method.		
KM	Signal sequence trap method; SST method; immunisation; inhibition;		
KM	infection; allergy; cancer; regulation; tissue formation; tissue repair		
KM	actin activity; inhibit activity; chemokine activity;		
KM	cytokine activity; blood coagulation regulation; agonist; antagonist;		
KM	metabolic disorder; hormonal disorder; immune disorder;		
KM	severe combined immunodeficiency; SCID; AIDS; thrombosis; cancer;		
KM	wound; ss.		
OS	Homo sapiens.		

XX	MO9918126-A1.
XX	
PD	15-APR-1999.
PF	06-OCT-1998; 98WO-JP04514.
PR	07-OCT-1997; 97JP-0274674.
PA	(ONOX) ONO PHARM CO LTD.
PI	Fukushima D, Shlbayama S, Tada H;
DR	WPI: 1999-277254/23.
DR	P-PSTDB: AAY02371.
PT	Polypeptides identified by the signal sequence trap method from a
PT	human CDNA library
XX	
XX	Claim 5; Page 139-140; 281pp; Japanese.
XX	
CC	AAV53694-X35747 represent cDNA sequences that encode novel polypeptides
CC	AAV02358-84) which are identified from a human placental cDNA library
CC	by the signal sequence trap (SST) method. The polypeptides have a
CC	broad range of physiological activity, including immunisation against
CC	and inhibition of infections, allergies and cancer; regulation of tissue
CC	formation and repair; activin/inhibin activity; chemokine/cytokine
CC	activity; blood coagulation regulation; and receptor/ligand agonist
CC	or antagonist activity. The polypeptides can be used for prevention
CC	and treatment of disorders including infections by bacteria, yeasts and
CC	viruses (including HIV) and protozoa; metabolic and hormonal disorders;
CC	immune disorders (including severe combined immunodeficiency (SCID)
CC	and AIDS; Chromoblasts; cancer; and traumatic or surgical wounds.
XX	
SQ	Sequence 1422 BP: 538 A; 213 C; 332 G; 339 T; 0 other;
Alignment Scores:	
Pred. No.:	0 231 Length: 1422
Score:	111.00 Matches: 95
Percent Similarity:	37.42% Conservative: 79
Best Local Similarity:	20.43% Mismatches: 165
Query Match:	4.71% Indels: 126
DB:	Gaps: 26
US-09-807-459-2 (1-458) x AAV35720 (1-1422)	
OY	62 SerValSerAlaTyrMetSerArgCysAlaLysGlnAspCysLeuThrIleuGlnSerIeu 81
DB	181 ACCATTAATAATGCTGTAGACAGTTTATGATGAGTGCGCAC-----AGCTTTGATGAGATG 234
OY	82 LysTyrProLeuGluAlaIleTytyGlnProLeuThrIleuProaspProtyrGlnIleuGlu 101
DB	235 AATGCAGAGCTGCAKGTCAAACCTGAAGGATTATTTAATGTGGATGCTTTAAAGTCGAAA 294
OY	102 -----AlaIaIaPheIleuPheIlys 108
DB	295 TCATTAGAAGCAAAAAACAGAGCATTTGATGAACAGATTCACAAGATTGGAAACAAGAAG 354
OY	109 GluSerAspAlaAsnProAlaAsnSerThyGluLys----- 120
DB	355 GAAAAAACCGAATCGTCTAGAGCTGTTGAGAAAACCTGAAGGCTTCCTTACAAGAGAT 414
OY	121 -----ArgPheTrpMetArgPheArgArgGlyLysAsnHisSerTyrPheHisAspLeu 138
DB	415 GTTCAAAAGTTCAGGCACTACATGACGAATTTGGAGAGTCTCATTCAGGCATTTCTTGACCAG 474
OY	139 ValPheAsnLeuLeuGluLysAsnValThrArg-----Asp 150
DB	475 AAATTAATGTGCTCAATAGAGAAATTGCTAGACTAGACTAGAACTGAAACATATAAA 534
OY	151 AlaAspAlaThrAspIleGluAsnPheAlaSerArgTyrIleuTyrMetaIaThrIleuTyr 170
DB	535 CAGGCAACACTCGCATTCAGATATCATTCATTGACAAACCAAGATGAC----- 579


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DB 1416 GATAGAGAAATATAGAGATCATGTCAGAGATCTTCGGAAT----- 1460
QY 429 ThrValAspPheIleuAsnLysGluIleArgAsp-----ProSerLysAlaLeuIleArg 446
DB 1461 -----ATTAAAGAGATTAGATAGATGAGAAAGAAAGCTACTCTAATT 1505
QY 447 LysValSerThrGlu 451
DB 1506 AAGCTCTCTGAGAA 1520
RESULT 30
AAAT73285
ID AAT73285 standard; cDNA; 3278 BP.
XX
AC AAT73285;
XX
DT 12-SEP-1997 (first entry)
DE
DE K. lactis origin of replication complex protein 1 gene.
XX
XX Origin of replication complex; ORC; yeast; Kluyveromyces fragilis;
XX chromatinography; peptide sequencing; primer; amplification; PCR; genome;
XX polymerase chain reaction; open reading frame; cell growth; cancer;
XX infection; inflammation; hypersensitivity; ds.
XX
OS Kluyveromyces fragilis.
XX
FH Key Location/Qualifiers
FT CDS 395..3052
FT /tag= a
FT /product= ORC 1 protein
XX
XX US5614618-A.
XX
XX 25-MAR-1997.
XX
XX 16-DEC-1993; 9305-0168479.
XX
XX 07-JUN-1995; 95US-0484106.
XX
XX 16-DEC-1993; 93US-0168479.
XX
XX (COLD-) COLD SPRING HARBOR LAB.
XX
XX (REGC) UNIV CALIFORNIA.
XX
XX Bell SP, Foss M, Gavin K, Herskowitz I, Hidaka M,
XX Kobayashi R, Laurensen P, Li J, McNally FJ, Rine J;
XX Stillman RW;
XX
XX WPI; 1997-201534/18.
XX
XX P-PSDB; AAW22230.
XX
XX Nucleic acids encoding origin of replication complex proteins - used
XX for screening for lead cnds. for therapy or diagnosis of disease
XX associated with undesirable cell growth
XX
XX Claim 4; Column 59-62; 53pp; English.
XX
XX This is the nucleotide sequence encoding the origin of replication
XX complex protein 1 (ORC1) from the yeast Kluyveromyces fragilis. The
XX sequence was isolated using primers based on amino acid sequence
XX conserved between the ORC1 and SIR3 proteins from Saccharomyces
XX cerevisiae. The amplified fragment was then used for low stringency
XX DNA hybridisation to obtain the K. lactis ORC1 gene sequence. The ORC
XX proteins (AAW2224-35) can be used to screen chemical libraries to
XX identify lead compounds useful in treatment and diagnosis of undesired
XX cell growth, e.g. cancer, infections, inflammation and hypersensitivity.
XX
XX Sequence 3278 BP; 1085 A; 572 C; 714 G; 902 T; 5 other;
XX
Alignment Scores: 0.705 Length: 3278
Pred. No.: 111.00 Matches: 98
Score:

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Percent Similarity: 35.50% Conservative: 82
Best Local Similarity: 19.33% Mismatches: 199
Query Match: 4.71% Indels: 128
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DB 1214 GAACCTATATCAGATTAATGATGATTCGATTAATATCAGAAAGTAAGAGACTTT 1273
QY 25 AlaAsnAlaIleMetIleAsnSerAsp-----MetSerAspTyrLeuSerAlaValSer 42
DB 1274 GCAACGACATCTCTTCGACAGATGAGAGATTGAAGATTAACAGCTCGAGAAAG 1333
QY 43 AspAsnPheAlaGluArgIleCysSerGluVal----- 53
DB 1334 CTTCGAATTTAGACACCTGCCAGAAAGGTCGATCTATTAAACGAGATATACCAT 1393
QY 54 ---ProLysGlySerAsnCysSerAlaSerValSerAlaIleMetSer-----Arg 69
DB 1394 TCACCACTTAATATCAGACATCCATTCAGCCATCAGACAGTTCATTCCTAGAAAG 1453
QY 70 CysAlaLysGlnAspCysLysLeuThrGlnSerLeuLysTyrProLeuGluAlaLysTyr 89
DB 1454 TTCCTTAAGATTAATATGATGACGCGCTAAAGGCAATATCTCTTCCAAACGGTAT 1513
QY 90 GlnProLeuThrLeuProAspProTyrGlnLeuGluAlaPheIleLeuPheLysGlu 109
DB 1514 AAGAAATCCGAAGATTCCTGCTGACATGACAT-----ATTTTCCAAAGCAT 1558
QY 110 SerAspAlaAsnProAlaAsnSerThrGluLysArgPheThrPheArgPheArgGly 129
DB 1559 AATAATGATTTGATATAGCGATTCATAGAGAGAGATTCGAACAGTTCTGCTAAAGC 1618
QY 130 LysAsnHisSerTyrPheHisAspLeuValPheAsnLeuGluLysAsnValThrArg 149
DB 1619 AAAATGAGACATATTTTTCAGAGAGAAAGCAATTCAGCAAGATATGCAAAAGAA 1678
QY 150 Asp---AlaAspAlaThrAspIleGluAsnPhe---AlaSerArgTyrLeuTyrMetAla 167
DB 1679 GAATTTGCAAAAGCTGCTGATTTGACAAATTTATTTCCGCGAGAGAAATGATTTGCA 1738
QY 168 ThrLeuTyrTyrLysThrTyrThrAsnValAspGluPheGlyAlaSerPheAsnLys 187
DB 1739 AGTATATACCTCTCATTACATGACGCAAT---GAAACGACGACCTACACCATTTTAC 1795
QY 188 LeuSerPheThrThrGlyLeuPheGlyTyrGlyIleLysArgAlaLeuLysGlnIle 207
DB 1796 ATTGCGGAGCGCCAGCGCTT-----GATAAACTTTGACGCTTCGAGAGTATGTT 1846
QY 208 ArgSerAsnLeuProLeuAspIleGlyThr-----GluHisSerValSerArgLeu 224
DB 1847 AAG-----GATTAATGACATCTCAGACCAAAAGAACTTCCAAAGTTTC 1891
QY 225 GlnHisIle-----ThrSerTyrLys----- 232
DB 1892 CAATACATTGAATCAATGCTTAAAGATTGTCAAAAGCAAGTATGATTAAGAGTCTT 1951
QY 233 1952 TGGCAAAAATATCTGAGAAAGCTTACATCTGACCTCCATGAGATCTGGAATTT 2011
QY 238 GlnIleProLeuLeuProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeu 257
DB 2012 TATTTTAACAAAGTTCAGCTAGCAAAAGCTCTATCGTTGCTTATTTGAGAGACTT 2071
QY 258 LeuAlaThrValAlaGlyTyrValAspThrProTyr---TyrLysLysTyr-----Tyr 274
DB 2072 GATGCATTAGTAGCAAGACCAAGATGATGATGATGATGATGATGATGATGATGAT 2131
QY 275 MetLysLeuLysAsnPheMetVal-----AsnArgValPheIleProThrLysLys 291

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Db 2132 TCAATGCGAAGCTTATGTGTAGCTGTGCGAAGACCTTAGATCTCCGAGCGCAT 2191
QY 292 Phepnealsnlys----- 295
Db 2192 CTGTGTAACAGATTTCTCCAGAAATGTTTACTAGAAATATGTCATGCTTACACG 2251
QY 296 -----guilelearg----- 298
Db 2252 CATGAGAGCTTAGACAAATCATCATTTGAGACTTAAATATTTGACGAATCTGTTTC 2311
QY 299 -----giuproserlysalaaleuylsaglulysvalserthrlyaspthrlyasphe 316
Db 2312 TATGTCGACCGGAGACAGGAGTGTACATGATCTCTCCGATAGTAGTACTATA--- 2368
QY 317 GLUsluylsllleglyglnlythrvalasphpheasnlsluileargasploser 336
Db 2369 GAAACTGATGAGAGAAAGACGAAAGACTTCTCTAAC-----TAT 2410
QY 337 Lysalaaleuylsaglulysvalserasnaspalalysasphegluasnlsllegly 356
Db 2411 AAACGACTTAAACTAGATTAATCCATGATGCCATGATTCATCAAGAAATTTGCT 2470
QY 357 Glnglythrvalasphpheileasnsluileargasploserlysalaaleuilearg 376
Db 2471 AGT-----gtcagtggtgagtgccgagagctttaaagtggtcAAAAAGA 2515
QY 377 Lysvalserthrnglyalaglaspheu-----Phegluasnlslile--- 390
Db 2516 GCGGTAGAAATTCGCGAAATGATTACTTAAAGAGCTTAGATAGCGAGCTACTCAAT 2575
QY 391 -----Glyglnlythrvalasphpheileasnsluile 402
Db 2576 TCCAAAAAAGATGACTAGTGCGCAATGTACAGAAATGAAGATTAACAGAGTGTAGAAAT 2635
QY 403 Argasploserlysalaaleu 409
Db 2636 AAGCATATTACCAAGGCATTA 2656

RESULT 31
AAT62358
ID AAT62358 standard; cDNA: 3278 BP.
XX
AC AAT62358;
XX
DT 23-JUL-1997 (first entry)
XX
DE Kluyveromyces lactis origin of replication complex ORC1 gene.
XX
KW Origin of replication complex; ORC; gene therapy; cancer;
XX
XX neoplasia; inflammation; hypersensitivity; ds.
XX
OS Kluyveromyces lactis.
XX
FH Key Location/Qualifiers
FT misc_difference 309
FT /*tag- a
FT /note= "base 309 is given as m in the
FT misc_difference 311
FT /*tag- b
FT /note= "base 311 is given as n in the
FT misc_difference 317
FT /*tag- c
FT /note= "base 317 is given as n in the
FT misc_difference 327
FT /*tag- d
FT /note= "base 327 is given as n in the
FT misc_difference 328
FT /*tag- e
FT /note= "base 328 is given as n in the
FT misc_difference 3143
FT /*tag- f
FT /note= "base 3143 is given as n in the

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FT CDS 395..1052 specification"
FT FT /*tag- f
FT PN W09640977-A1.
FT PD 19-DEC-1996.
FT PF 07-JUN-1996; 96WO-US09403.
FT PR 07-JUN-1995; 95US-0484105.
FT PA (COLD-) COLD SPRING HARBOR LAB.
FT PA (RESC ) UNIV CALIFORNIA.
FT PI Bell SP, Foss M, Herskowitz I, Kobayashi R, Laurensen P;
FT PI Li JT, McNally FJ, Rine J, Stillman BW;
FT DR WPI; 1997-052354/05.
FT DR P-PSDB; AAM14136.
FT XX Nucleic acid encoding origin of replication complex (ORC) protein -
FT PT useful to screen for lead pharmaceuticals capable of disrupting ORC
FT PT protein function, and inhibiting cell growth
FT PS Disclosure: Page 16-18; 57pp; English.
FT XX
CC Isolated cDNA clones (AAT62358-63) respectively encode origin of
CC replication (ORC) proteins (AAM14136-41) from Kluyveromyces lactis,
CC Schizosaccharomyces pombe, human (ORC1), Arabidopsis thaliana,
CC Caenorhabditis elegans and human (ORC2). The K. lactis ORC1 clone
CC was obtained by PCR amplification using primers based on Saccharomyces
CC cerevisiae ORC1 and SIR3 sequences. The isolated nucleic acids can
CC be utilised in the prodn. of ORC polypeptides, to design probes and
CC primers for the detection and amplification of ORC genes, and in
CC gene therapy appls., e.g. antisense oligonucleotides capable of
CC inhibiting the intracellular expression of a targeted ORC
CC transcript.
CC XX
SQ Sequence 3278 BP; 1085 A; 572 C; 714 G; 902 T; 5 other:

Alignment Scores:
Pred. No.: 0.705 Length: 3278
Score: 111.00 Matches: 98
Percent Similarity: 35.50% Conservative: 82
Best Local Similarity: 19.33% Mismatches: 199
Query Match: 4.71% Indels: 128
DB: 18 Gaps: 22

US-09-807-459-2 (1-458) x AAT62358 (1-3278)
QY 5 AspSerValGlyAspValThrlyThrleuLeuAlaSerGluSerValaspSerAla 24
DB 1214 GAAGCTATATACAGATTAATGCAATCGATTATGCAATATACGAAAGAGATT 1273
QY 25 AlaasnAlaTyMetileasnSerasp-----MetSerAspTyLeuSerAlaValser 42
DB 1274 GCAAAAGCATCTCTTCGACAGTGTGAAGAGTTTGAAGATTTACAGTCTGCAGAAAG 1333
QY 43 AspAsnPhelaIguArgIleCysSerGlnAl----- 53
DB 1334 CTTCGAATGTGTAACCTCCCAAGAAAGGTGAGATCTATTAAACCAATATACCAT 1393
QY 54 ---ProLysGlySerAsnCySerSerAlaSerAlaTyMetSer-----Arg 69
DB 1394 TCACCAAGTAAATACACAGATCCATTCAGCCATCGAGTTCATTCATCTCCAGAAAG 1453
QY 70 CysAlaLysGlnAspCysLeuThrleuGlnSerleuTyProleuAluAlaLysTyR 89
DB 1454 TTCCTTAAAGATTAATATATGCGCGCTAAAGAGCATATATCTCTTCCAAAGGTA 1513
QY 90 GlnProLeuThrleuProAspProTyrgInleuGlnAlaAlaPheIleuPheLysGlu 109
DB 90

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Alignment Scores:

Pred. NO.:	0.391	Length:	1777
Score:	110.00	Matches:	97
Percent Similarity:	35.04%	Conservative:	67
Best Local Similarity:	20.73%	Mismatches:	170
Query Match:	4.66%	Indels:	134
DB:	11	Gaps:	24

US-09-807-459-2 (1-458) x AA006842 (1-1777)

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OY 23 SerAlaAlaSerAlaTyrMetIleAsnSerAspMetSerAspTyr----- 37
Db 332 AACUCUGAAGUAGAAUUGUUUACAGAGAUUUUUAUCUGACUAGUAGAGAGCGAG 391
OY 38 ---LeuSerAlaValSerAspAsnPhaIleArgIleCysSerGlnValProIysGly 56
Db 392 CAUUGGAGCUCAGUGUACA-----UUGCAAGAGAUUC---GAAAUUUUUCCAAAGAA 442
OY 57 Ser-----AsnCysSerAlaSerValSerAlaTyrMetSerArgCysAla 71
Db 443 AGCUCUAGUGCCCAACACACACAAACAAAGAGUAGCGGCAUCCGCAUGCGGGG 502
OY 72 LysGlnAsp-----CysIleuThrLeuGlnSerLeuIysTyrProIeu 85
Db 503 AAAAGCAGUUUUUACGAGAAUUUGCUAUGGCGAGAGAGAGGCGUACAACCA--- 559
OY 86 GluAlaIysTyrGlnProIeuThrIleuProAspProTyrGlnLeuGlnAlaIaPhelle 105
Db 560 -----AACGUGAAAAUUUCUUUUGUG 580
OY 106 LeuPheLysGluSerAspAla-----AspProAlaAsnSerThr 118
Db 581 AACAGAAAGGAAAGAAAGUCCUUUACUGUGGUGGUUUUACACACCGGCUAACAGUAG 640
OY 119 GluLysArgPheTyrMetIleArgIleGlyLysAsnHisSerTyrPheHisAspIeu 138
Db 641 GAUCCAACAG-----AAUUAUCUACAGAAUAAU---GCUUAUGUCUCUGUAGUG 688
OY 139 ValPheAsnLeuGlnLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsn 158
Db 689 ACUUCAAAUUUUAAACAGGAGAUUUUACCCCGGAAUAGCGAAAGACCCAAAUUAGAGAGU 748
OY 159 PheAlaSerArgTyrLeuTyrMetAlaThrIleuTyrTyrTyrThrAsnValAsp 178
Db 749 CAAGCUGGAGAGAUACUAAUUAUCUGACCUUUGCUAAAAACCGGAGACACAAUAAUUAU 808
OY 179 GluPheGlyAlaSerPhePhe-----AsnLysLeuSerPheThrThrGlyLeuPhe 195
Db 809 GAGGCAAAUUGAAAUUCUAAUACGACCAAGUUAUGUUUGCAGUCAGUGAGGCG---UUU 865
OY 196 GlyTyrGlyIleLysArgAlaLeuLysGlnIleIleArgSerAsnLeuProIeuAspIle 215
Db 866 GGGUCCGCG-----AUCACUCCUCAAAGCAACGCAUCAAUAGCAGUGAG 904
OY 216 GlyThrGlnHisSerValSerArgLeuGlnHisIleThrSerSer-----TyrLysAsp 233
Db 905 UGUAAACAGAAUGUCAAACACCCUUGGAGAGUAUAAACAGCAGUCUCCUUUCCAGAAU 964
OY 234 TyrMetAspThrGlnIlePheAlaLeuProLysPheAlaLysArgPheSerLeuMetVal 253
Db 965 AUAACACCAAGUCACAUAUUGAGAGUGCCAAAUUACGUCAGAGAGGCCAAAUUUGAGAGUG 1024
OY 254 Val-----GlnArgLeuLeuAlaThrValAla 262
Db 1025 GUUACAGAGCUAAGAGACAUUCCGCUAUUCCAGGAGGCUUUUUUGAGCCAUUUGCC 1084
OY 263 GlyTyrValAspThrProTyr-----TyrLysTyrTyrTyr----- 274
Db 1085 GGUUUUUUUAAGAGGGAGGAGCUGAGAAUAGAGAGUGUAGCGUUUUAUCUACUACAG 1144
OY 275 -----MetLysLeuLysAsnPh 280

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```

Db 1145 AAUAGACAGGAGUACAGGCUUAGCAGCGAGUACAAAAAGCACACAAAAUUGCCAUUAAACGG 1204
OY 281 MetValAsnArgVal-----PheIleProThrLys 290
Db 1205 AUUACAAACAAAGGUGUAAUCUUGUACGAGAAAUAGACAUUCCAUUCCACAGCUGUGGU 1264
OY 291 LysPhePheAsnLysGluIleArgGluProSerLysAlaLeuLysGluLysValSerThr 310
Db 1265 AAAGAUAUUCACAA---UUAGAAAAAGAGUAGAAAUUUUAUAAAAAGUUGAU 1321
OY 311 AspThrLysAsp-----LeuPheGluAsnLysIle 320
Db 1322 GGAUUUUCUGACAUUUGGACAUUAUAGCAGAUUGCUUAGUUCUACUGGAAAAU----- 1375
OY 321 GlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspProSerLysAlaLeuLys 340
Db 1376 ---GAAAGAGCUCUGAGUUUCCAUAGACUCAAUUGUCAAACAUUCUAGUAGAAAGUAAA 1432
OY 341 GluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThrVal 360
Db 1433 AGCCAAUUAAGAAUAUAUAGCAAAGAA-----AUCGAAUUGGAGUUGUU 1477
OY 361 AspPhe-----IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 377
Db 1478 GAGUUCUACCAAGAGUGUAGCAUAGAA-----UGCAUUGGAAAGU 1516
OY 378 ValSerThrGlyAlaGluAspLeu-----PheGluAsnLysIleGlyGlnGly 393
Db 1517 GUAAGAAUUGGACUUAUAGAUUUAUCCCAAUAUUAUUCAGAAAGAGUCAAAGUAAACAGGAA 1576
OY 394 ThrValAspPheIleAsnAsnGlu 401
Db 1577 AAGGUAGAGAGAGUAGAAUUGGAA 1600

RESULT 33
AA574418
ID AA574418 standard; cDNA; 1816 BP.
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AC AA574418;
XX
DE 13-FEB-2002 (first entry)
XX
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XX DNA encoding novel human diagnostic protein #10222.
XX
XX Human; Chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
XX Homo sapiens.
XX
XX WO200175067-A2.
XX
XX 11-OCT-2001.
XX
XX 30-MAR-2001; 2001WO-US08631.
XX
XX 31-MAR-2000; 2000US-0540217.
XX
XX 23-AUG-2000; 2000US-0649167.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Drmanac RT, Liu C, Tang YT;
XX
XX WPI: 2001-639362/73.
XX
XX P-PSDB: ABG10231.
XX
XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity -
XX
XX Claim 1; SEQ ID No 10222; 103pp; English.
XX
XX The invention relates to isolated polynucleotide (I) and

```

CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations in
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at [ftp.wipo.int/pub/published_pct_sequences](http://wipo.int/pub/published_pct_sequences).

XX Sequence 1816 BP; 726 A; 380 C; 328 G; 382 T; 0 other;

Alignment Scores:
Pred. NO.: 0.45 Length: 1816
Score: 109.50 Matches: 76
Percent Similarity: 37.99% Conservative: 41
Best Local Similarity: 24.68% Mismatches: 113
Query Match: 4.64% Indels: 78
DB: 23 Gaps: 16

US-09-807-459-2 (1-458) x AAS74418 (1-1816)

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Db 721 ACTTCTTACAGAAATTTGAAAAAATCTTAAAGTTCAATGAAACCAAAAAAGACC 780
QY 203 LeuLysGlnIleIleArgSerAsnLeu--ProLeuAspIleGlyThrGlnHisSerVal 222
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Db 781 TGC-----ATTGCCAAGTCAATCTTAAGCCAAAGACGAGCATCTGCTA 834
QY 222 eATGLeuGlnHisIleThrSerGlyLysAspTyrMetAspThrGlnIleProAla 242
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Db 835 CCGGACTTCAAACTATATACAGGCTACAGTAACT-----CAAAACAGCATGTACTGG 888
QY 242 euProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuAlaThrValA 262
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Db 889 TACCAAAA-----CAGGATATAGACCAATGGAACAGACAGCCCTCAGAAATATG 942
QY 262 LaglyTyr-----Val-AspThrPro-----Trp----- 269
|||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 943 CCGCATATCTACAACTATCTGATCTTCGACAAACCTGACCAAAACAGCAATGGGAAAG 1002
QY 270 -----TyrLysLysTyrTyr-----MetLys 276
:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 1003 GATTCCCTATTAAATTAATGGTACTGCGAAAACTGCTAGCCATATGTAGAAAGCTGAAA 1062
QY 277 LeuLysAsnPhmetValAsnArgValPheIleProThrLysLysPheAsnLysGlu 296
|||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 1063 CTAATCCCTTCTTACACCTTATACAAAAATTAATTAACATGATGATTAAGACTTAAT 1122
QY 297 IleArg---GluProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeu 315
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QY 316 PheLysAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspPro 335
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Db 1180 -----ATAGGCGATGGCAAG---GACTTCATGTCTTAA-----ACA 1212
QY 336 SerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGluAsnLys--- 354
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Db 1213 CCAAAAGCAATGGCAACAAAGACAAATTCACAAATGGGATCTAATTAACCAAGAGGC 1272

QY 355 -----IleGlyGlnGlyThrValAspPheIleAsnGlnIleIleArgAspProSerLys 372
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Db 1273 TTTCGACAGCAAAAGAAACATACCATCTCAGAGTGAAAC-----AGGCAACCTCAAAA 1323
QY 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGln 392
|||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 1324 -----TGGAGAAATAATTTCCGCAAC----- 1344
QY 393 GlyThrValAspPheIleAsnGlnIleIleArgAspProSerLysAlaLeuIleArgLys 412
1345 -----TACTCATCTGACAAAGGCTTAATATCCAGA 1374
QY 413 ValTyrThrGlnAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe 432
|||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 1375 AACTACATTAAGAACTCAACCAATTTACAGAAAA-----ACAATGGGCCCC 1422
QY 433 IleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysValSerThrGlnAla 452
|||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 1423 ATCAAAAAGTGGCGAGAGGATATGAAACAGCATCTTCAAAAAGAACACATTATGACGCC 1482
QY 453 AspAsnLeuLeuGluLys 458
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Db 1483 AAAAAACATGAAAAAA 1500
RESULT 34
AAS69641
ID AAS69641 standard; cDNA; 2277 BP.
XX AAS69641:
AC AAS69641:
XX 13-FEB-2002 (first entry)
DT
XX
DE DNA encoding novel human diagnostic protein #5445.
KW Human: chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN MO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI: 2001-639362/73.
DR P-PSDB; ABG05454.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX
PS Claim 1; SEQ ID No 5445; 103pp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating

CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 2277 BP: 878 A; 483 C; 473 G; 443 T; 0 other:

Alignment Scores:
 Pred. No.: 0.609 Length: 2277
 Score: 109.50 Matches: 73
 Percent Similarity: 37.21% Conservative: 39
 Best Local Similarity: 24.25% Mismatches: 107
 Query Match: 4.64% Indels: 82
 Gaps: 15

US-09-807-459-2 (1-458) x AAS69641 (1-2277)

QY 183 SerpPheasnLysLeuSerPheThrGlyLeuPheGlyTrpGlyIleLysArgala 202
 Db 868 ACTTCTCCGAGATGGAATAAACTTAAAGTTCATATGACCAAAAGAGGCC 927
 QY 203 LeuLysGlnIleIleArgSerAsnLeu--ProLeuAspIleGlyThrGlnHisSerValS 222
 Db 928 CTC-----ATCGCCAGCATCTCTGAGCCAAAGAACCCAGAGACATCAGCTCA 981
 QY 222 eArgLeuGlnHisIleThrSerSerTrpLysAspTrpMetAspThrGlnIleProAlaL 242
 Db 982 CCTGACTTCGAGACTTACTACAGGCTACAGTAC-----CAAACAGCTGGTACTG 1035
 QY 242 euProlLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuAlaThrVal 262
 Db 1036 TACCAAAA-----CAGAGATATAGACCAATGAGACAGACAGCCCTCAGAAATATAG 1089
 QY 262 IagLYTyr-----Val-AspThrPro----- 268
 Db 1090 CCGCATATCTACACTATCTGATCTTTGACAAACCTCACAAAACAGCAATGGGGAAG 1149
 QY 269 -----Tyr---TyrLysLysTrpTyrMetLysLeuLysAsnPhemet 281
 Db 1150 GATTCCTATTTAATTAATGCGCTGGGAAAAATGCTAACATATGTGAAGAAGCTGAAA 1209
 QY 282 ValAsnArgValPheIleProThrLysLysPheAsnLysGluIleArgGlu----- 299
 Db 1210 CTGATCCCTTCCCTTACACCTTATCAAAAATTAATTCAGATGATTAAGACTTACAT 1269
 QY 300 -----ProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeu 315
 Db 1270 GTTAGAGCTAAACCATTAATAAACCCTAGAGAAAACCTAGGCAATACCATTTAGAGC 1326
 QY 316 PheGlnAsnLysIleGlyGlnIleThrValAspPheAsnLysGluIleArgAspPro 335
 Db 1327 -----ATAGGCATGGCAAG---GACTCATGTCTTAAA-----CCA 1359
 QY 336 SerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGlnAsnLys 354
 Db 1360 CCAAAAGCAGCGCAGCAAAAGCCAAATATGCAAAATGGAGTCTAATTAACATAAAGAAC 1419
 QY 355 -----IleGlyGlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLys 372
 Db 1420 TTCTGCACAGCAAGAAAGAACTACCATCAGAGTGAAC-----AGGCAACCTCAAAA 1470
 QY 373 AlaLeuIleArgLysValSerThrGlyAlaGlnAspLeuPheGlnAsnLysIleGlyGln 392
 Db 1471 -----TGGGAGAAATTTTACAAAC----- 1491
 QY 393 GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 412

Db 1492 -----TACTCATCTGACAAAGGCGTAATATCCAGA 1521
 QY 413 ValTyrThrGluAlaAspAspLeuPheGlnAsnLysIleGlyGlnIleThrValAspPhe 432
 Db 1522 ACTTACATGAACTCCAGCAAAATTTACAGAAAAA----- 1557
 QY 433 IleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysValSerThrGln 451
 Db 1558 ACAAAACACCCCATCAAAAAGTGGGGAAGCATGACAGACATCTTCAAAAAGA 1614

RESULT 35

AAS71279 standard; cDNA; 2277 BP.
 ID AAS71279
 AC AAS71279;
 XX 13-FEB-2002 (first entry)
 DT 13-FEB-2002 (first entry)
 DE DNA encoding novel human diagnostic protein #7083.
 XX
 KM Human: chromosome mapping; gene mapping; gene therapy; forensic;
 XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001WO-US06831.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 PA (HSE-) HYSEQ INC.
 XX
 PI Drmanac RT, Liu C, Tang YT;
 XX
 DR WPI: 2001-639362/73.
 DR P-PSDB: ABO7092.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity
 XX
 PS Claim 1: SEQ ID No 7083; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 2277 BP: 878 A; 483 C; 473 G; 443 T; 0 other:
 Alignment Scores:

QY 269 -----TTP---TyrLysLysTrpTyrMetLysLysAsnPhmet 281
DB 1150 GATTCCCTATTATAAATGCTGGGAAAAAATGCTAACCAATATGTAAGAAACCTGAAA 1209
QY 282 ValAsnArgValPheIleProThrLysLysPhePheAsnLysGluIleArgGlu----- 299
DB 1210 CTGATCCCTCTTCCATACCTTATACAAAATTAATTCAGATGATGATTAAGACTTACAT 1269
QY 300 -----ProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeu 315
DB 1270 GTTAGAGCTAAACCATTAACCTTGAAGAAAAGAACTAGCAATACCATTCAGAC--- 1336
QY 316 PheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspPro 335
DB 1327 -----ATAGGCATGGGCAAG---GACTTCATGCTTAA-----CCA 1359
QY 336 SerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGluAsnLys--- 354
DB 1360 CCAAAACCAACGGCAGCAAAAGCCAAATTCACAAATGGATCTTAATTAACCTAAAGAAC 1419
QY 355 -----IleGlyGlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLys 372
DB 1420 TTCTGCACAGCAAAAGAACTACCATCAGACTGAC-----AGCAACCTACAAA 1470
QY 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGln 392
DB 1471 -----TGGGAGAAATTTTCACACAC----- 1491
QY 393 GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 412
DB 1492 -----TACTCATCTGACAAAGGGCTTAATTCACAGA 1521
QY 413 ValTyrThrGluAlaAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe 432
DB 1522 ATCTACATGACATCCAGCAATTTACAGAAAA----- 1557
QY 433 IleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysValSerThrGlu 451
DB 1558 ACAAAACCAACCCATCAAAAGTGGCGACAGCATGAACACACTCTCAAAAGAA 1614
RESULT 38
AA574599
ID AA574599 standard; cDNA; 2277 BP.
XX
AC AA574599;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #10403.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001MO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
XX
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI; 2001-639362/73.
XX
DR P-PSDB; ABG10412.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in

PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
PS Claim 1; SEQ ID No 10403; 103bp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AA564197-AA594564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pcl_sequences.
XX
SQ Sequence 2277 BP; 878 A; 483 C; 473 G; 443 T; 0 other;
Alignment Scores:
Pred. No.: 0.609 Length: 2277
Score: 109.50 Matches: 73
Percent Similarity: 37.21% Conservative: 39
Best local Similarity: 24.25% Mismatches: 107
Query Match: 4.64% Indels: 82
DB: Gaps: 15
US-09-807-459-2 (1-458) x AA574599 (1-2277)
QY 183 SerPhePheAsnLysLeuSerPheThrGlyLeuPheGlyTyrPglLysArgAla 202
DB 868 ACTTCTTCCAGAAATTTGGAAAAAACTTAAAGTTCAATATGAAACCAAAAGAACCC 927
QY 203 LeuLysGluIleIleArgSerAsnLeu--ProLeuAspIleGlyThrGluHisSerValS 222
DB 928 CTC-----ATCGCCAGTCAATCCTGAGCCAAAGAACCAAGCAGATCAGCCTA 981
QY 222 eraTgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIleProAlaL 242
DB 982 CCGTACTTCGAACATATCTACAGGCTACAGTAC-----CAAAACAGCCTGGTACTGG 1035
QY 242 eupProLysPheAlaLysArgPheSerLeuMetValIleGlnArgLeuLeuIleThrValA 262
DB 1036 TACCAAAA-----CAGAGATATAGACCAATGGAACAGACAGCCCTCAGAAATATAG 1089
QY 262 laGlyTyr-----Val-AspThrPro----- 268
DB 1090 CCGCATATCTACAACTATCTGATCTTGACAAACCTCAGAAAAACCAAGCAATGGGAAAG 1149
QY 269 -----TTP---TyrLysLysTrpTyrMetLysLysAsnPhmet 281
DB 1150 GATTCCCTATTATAAATGCTGGGAAAAAATGCTAACCAATATGTAAGAAACCTGAAA 1209
QY 282 ValAsnArgValPheIleProThrLysLysPhePheAsnLysGluIleArgGlu----- 299
DB 1210 CTGATCCCTCTTCCATACCTTATACAAAATTAATTCAGATGATGATTAAGACTTACAT 1269
QY 300 -----ProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeu 315
DB 1270 GTTAGAGCTAAACCATTAACCTTGAAGAAAAGAACTAGCAATACCATTCAGAC--- 1336
QY 316 PheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspPro 335

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Db 1327 -----ATAGGCAATGGGCAAG---GACCTTCATGCTCTAA-----CCA 1359
Qy 336 SerLysAlaLeuLysGluValSerAsnAspAlaLysAspLeuPheGluAsnLys--- 354
Db 1360 CCAAAAGCAACGGCAGCAACCAAAATTTGACAAATGGGATCTTAATTAACCTAAAGAAC 1419
Qy 355 -----IlleGlyGlnGlyThrValAlaAspPheIleAsnAsnGluIleArgAspProSerLys 372
Db 1420 TTCTGCACAGCAAAAGAAATACCATCAGAGTGAC-----AGGCACTTACAAAA 1470
Qy 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGln 392
Db 1471 -----TGGAGAAATTTTTCACACC----- 1491
Qy 393 GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 412
Db 1492 -----TACTCATCTGACAAAGCGCTAAATATCCAGA 1521
Qy 413 ValTyrThrGluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe 432
Db 1522 ATCTACAAATGAACTCCAGCAAAATTTACAAAGAAAA----- 1557
Qy 433 IleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysValSerThrGln 451
Db 1558 ACAAAACAAACCCCATCAAAAAGTGGCGAAGACATGACACACATCTCAAAAAGAA 1614

RESULT 39
AAS79126
ID AAS79126 standard; cDNA; 2277 BP.
XX
AC AAS79126;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #14930.
XX
KW Human; Chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
XX
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI: 2001-639362/73.
XX
DR P-PSDB; ABG14939.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity.
XX
PS Claim 1; SEQ ID No 14930; 103pp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or

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CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AS64197-AS64554 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 2277 BP; 878 A; 483 C; 473 G; 443 T; 0 other;

Alignment Scores:
Pred. No.: 0.609 Length: 2277
Score: 109.50 Matches: 73
Percent Similarity: 37.21% Conservative: 39
Best Local Similarity: 24.25% Mismatches: 107
Query Match: 4.64% Gaps: 82
DB: 23 Indels: 15

US-09-807-459-2 (1-458) x AAS79126 (1-2277)
Qy 183 SerPhePheAsnLysLeuSerPheThrGlyLeuPheGlyTyrGlyIleLysArgAla 202
Db 868 ACTTCTCTCCAGAAATTTGAAAAAACTACTTTAAAGTCTATATGAAACCAAAAAGAGCC 927
Qy 203 LeuLysGlnIleIleArgSerAsnLeu--ProLeuAspIleGlyThrGluIleSerVal 222
Db 928 CTC-----ATGCCCAAGTCAATCTGAGCCAAAGAACCAACCAAGACATCAGCGTA 981
Qy 222 eraArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIleProAla 242
Db 982 CCTGACTTGCAGTAACTATCTCAAGAGCTACAGTAAAC-----CAAAACAGCTGACTCG 1035
Qy 242 eupProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThrValA 262
Db 1036 TACCAAAA-----CAGAGTATAGACCAATGGAACAGAACAGACCCCTCAGAAATATG 1089
Qy 262 IagLysTyr-----Val-AspThrPro----- 268
Db 1090 CCGCATATCTACAACTATCGATCTTGCAAAACCTCACAAAACCAACATGGGGAAG 1149
Qy 269 -----Tyr--TyrLysLysTyrTyrIleLysLeuLysAsnProMet 281
Db 1150 GATTCCCTATTATTAATAATGTCTGCGAATAATGGCTACCATATGTAGAAAGCTGA 1209
Qy 282 ValAsnArgValPheIleProThrLysLysPhePheAsnLysGluIleArgGlu 299
Db 1210 CTGGATCCCTCTCTACACCTTATACAAAATAATTAATTAAGATTAAGACTTACAT 1269
Qy 300 -----ProSerLysAlaLeuLysGluValSerThrAspThrLysAspLeu 315
Db 1270 GTTAGAGCTTAACCATTAATAAACCTTAGAAGAAAACCTAGCATTCACAGC--- 1326
Qy 316 PheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspPro 335
Db 1327 -----ATAGGCAATGGGCAAG---GACCTTCATGCTCTAA-----CCA 1359
Qy 336 SerLysAlaLeuLysGluValSerAsnAspAlaLysAspLeuPheGluAsnLys--- 354
Db 1360 CCAAAAGCAACGGCAGCAACCAAAATTTGACAAATGGGATCTTAATTAACCTAAAGAAC 1419
Qy 355 -----IlleGlyGlnGlyThrValAlaAspPheIleAsnAsnGluIleArgAspProSerLys 372
Db 1420 TTCTGCACAGCAAAAGAAATACCATCAGAGTGAC-----AGGCACTTACAAAA 1470
Qy 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGln 392
Db 1471 -----TGGAGAAATTTTTCACACC----- 1491

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QY 393 GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 412
Db 1492 -----TACTCATCTGCACAAAGGCTAATATCCAGA 1521
QY 413 ValTyrThrGluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe 432
Db 1522 ATCTACATGAACTCCACCAATTTTACAAAGAAAA----- 1557
QY 433 IleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysValSerThrGlu 451
Db 1558 ACAAAACAAACCCATCAAAAGTGGCGAAGACATGACAGCACTTCTCAAAAGAA 1614
RESULT 40
AAS84093
ID AAS84093 standard; cDNA; 2394 BP.
AC AAS84093;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #19897.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HXSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI: 2001-639362/73.
DR P-PSDB: ABG19906.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX
PS Claim 1; SEQ ID No 19897; 103bp; English.
XX
CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridization probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS84197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX

SQ Sequence 2394 BP; 935 A; 504 C; 490 G; 465 T; 0 other;
Alignment Scores:
Pred. No.: 0.651 Length: 2394
Score: 109.50 Matches: 73
Percent Similarity: 37.21% Conservative: 39
Best Local Similarity: 24.25% Mismatches: 107
Query Match: 4.64% Indels: 82
DB: 23 Gaps: 15
US-09-807-459-2 (1-458) x AAS84093 (1-2394)
QY 183 SerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrPglYIleLysArgAla 202
Db ACTTCTTCCCGAATTCGAAATTCCTTAAAGTTCAATGACCAACCAAAAGAGACC 1044
QY 203 LeuLysGlnIleIleArgSerAsnLeu--ProLeuAspIleGlyThrGlnLysSerValS 222
Db 1045 CTC-----ATCGCCAGCATCTGAGCCAAAGAACCAAGCAAGCATCAGCTA 1098
QY 222 erArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIleProAlaL 242
Db 1099 CTGACCTTCGAACTTACTACAGGCTACAGTAAAC-----CAAAACAGCCTGACTGG 1152
QY 242 euProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuAlaThrValA 262
Db 1153 TACCAAAA-----CAGAGATATAGCAATGGAACAGACAGACCCCTCAGAAATATG 1206
QY 262 laglyTyr-----Val-AspThrPro----- 268
Db 1207 CCGCATATCTACAACTATCTGATCTTGCACAAACCTCAAAACAGCAATGGGGAAG 1266
QY 269 -----Trp---TyrLysLysTyrTyrMetLysLeuLysAsnPheMet 281
Db 1267 GATTCCTTATTAAATGAATGGTGGGGAAGAAATGCGTAAACCATATGTGAAGAGCTGAAA 1326
QY 282 ValAsnArgValPheIleProThrLysLysPhePheAsnLysGluIleArgGlu----- 299
Db 1327 CTGGATCCCTTCCTTACACCTTATACAAATAATTAACTGATGGATTAAGACTTACAT 1386
QY 300 -----ProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeu 315
Db 1387 GTTAGAGCTAAACCATTAATAACCTTAGACAGAAACATGACCATACATTACAGAC--- 1443
QY 316 PheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspPro 335
Db 1444 -----ATAGGCATGGCAAG---GACTTCATGCTTAAA-----CCA 1476
QY 336 SerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGluAsnLys--- 354
Db 1477 CCAAAACGACAGGCGACGAAAGCCAAATTTGACAAATGGATCTTAATTAACTAAAGAAC 1536
QY 355 -----IleGlyGlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLys 372
Db 1537 TTTCGACACAGCAAAAGAAATCATCATCAGAGTAAAC-----AGGCACACCTACAAA 1587
QY 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGln 392
Db 1588 -----TGGGAGAAATAATTTTCACAAAC----- 1608
QY 393 GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 412
Db 1609 -----TACTCATCTGCACAAAGGCTAATATCCAGA 1638
QY 413 ValTyrThrGluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe 432
Db 1639 ATCTACATGAACTCCACCAATTTTACAAAGAAAA----- 1674
QY 433 IleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysValSerThrGlu 451
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Job time : 1278 secs

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